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UTILITY PATENT APPLICATION TRANSMITTAL
(new nonprovisional applications under 37 CFR 1.53(b))

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APPLICATION ELEMENTS

1. ☐ Fee Transmittal Form
(original and duplicate)
2. ☒ Specification [Total Pages 95]
3. ☐ Drawings [Total Sheets]
4. Oath or Declaration [Total Pages]
 - a. ☐ Newly executed (original or copy)
☐ New (unexecuted)
 - b. ☐ Copy from a prior application
(for continuation/divisional with
Box 17 completed)
 - i. ☐ DELETION OF INVENTOR(s)
Signed statement attached
deleting inventor(s) named
in prior application.
5. ☐ Incorporation By Reference
(useable if Box 4b is marked)

The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 4b, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein.

6. ☐ Microfiche Computer Program (Appendix)
7. ☐ Nucleotide and/or Amino Acid Sequence Submission
(if applicable, all necessary)
- a. ☐ Computer Readable Copy
- b. ☐ Paper Copy (identical to computer copy)
- c. ☐ Statement verifying identity of above copies

ACCOMPANYING APPLICATION PARTS

8. ☐ Assignment Papers (cover sheet & document(s))
9. ☐ 37 CFR 3.73(b) Statement ☐ Power of Attorney
10. ☐ English Translation Document (if applicable)
11. ☐ IDS with PTO-1449 ☐ Copies of IDS Citations
12. ☐ Preliminary Amendment
13. ☒ Return Receipt Postcard
14. ☐ Small Entity Statement(s)
☐ Statement filed in prior application; status still proper and desired
15. ☐ Certified Copy of Priority Document(s) if foreign priority is claimed
16. ☐ Other: _____

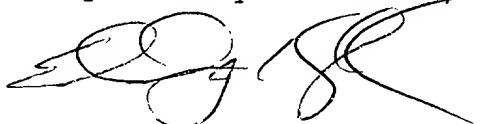
IF A CONTINUING APPLICATION, CHECK APPROPRIATE BOX AND SUPPLY THE REQUISITE INFORMATION

17. ☐ Continuation ☐ Divisional ☐ Continuation-in-Part of prior application No.: _____
- ☒ Complete Application
based on provisional Application No. 60/056,000; and
based on provisional Application No. unknown filed on
April 9, 1998 entitled PROCESS FOR SELECTIVE
DERIVATIZATION OF TAXANES.

CORRESPONDENCE ADDRESS

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Respectfully submitted,



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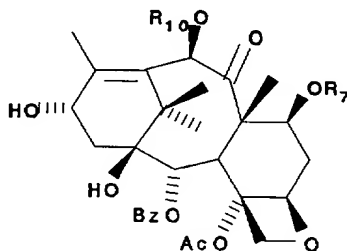
PROCESS FOR SELECTIVE DERIVATIZATION OF TAXANES

This invention was made with Government support under NIH Grant #CA 42031 awarded by the National Institutes of Health. The Government has certain rights
5 in the invention.

BACKGROUND OF THE INVENTION

The present invention is directed, in general, to a process for the preparation of taxol and other taxanes, and in particular, to such a process in which the C(7) or
10 C(10) hydroxyl group of a taxane is selectively derivatized.

10-DAB (1), which is extracted from the needles of *taxus baccata* L., the English yew, has become a key starting material in the production of taxol and
15 Taxotere, both of which are potent anticancer agents. Conversion of 10-DAB to taxol, Taxotere® and other taxanes having antitumor activity requires protection or derivatization of the C(7) and C(10) hydroxyl groups followed by esterification of the C(13) hydroxyl group to
20 attach an appropriate side chain at that position.



- 1 R₁₀ = R₇ = H
2 R₁₀ = H, R₇ = TES

Until now, strategies for the preparation of taxol
25 and taxol analogs were based upon the observation of

Senilh et al. (*C.R. Acad. Sci. Paris, II*, 1981, 293, 501) that the relative reactivity of the four hydroxyl groups of 10-DAB toward acetic anhydride in pyridine is C(7)-OH > C(10)-OH > C(13)-OH > C(1)-OH. Denis, et. al. reported (*J. Am. Chem. Soc.*, 1988, 110, 5917) selective silylation of the C(7) hydroxyl group of 10-DAB with triethylsilyl chloride in pyridine to give 7-triethylsilyl-10-deacetyl baccatin (III) (2) in 85% yield. Based upon these reports, in those processes in which differentiation of the C(7) and C(10) hydroxyl groups is required (e.g., preparation of taxol from 10-DAB), the C(7) hydroxyl group must be protected (or derivatized) before the C(10) hydroxyl group is protected or derivatized. For example, taxol may be prepared by treating 10-DAB with triethylsilyl chloride to protect the C(7) hydroxyl group, acetylating the C(10) hydroxyl group, attaching the side chain by esterification of the C(13) hydroxyl group, and, finally, removal of protecting groups.

It is known that taxanes having various substituents bonded to either the C(10) or the C(7) oxygens show anticancer activity. To provide for more efficient synthesis of these materials, it would be useful to have methods which permit more efficient and more highly selective protection or derivatization of the C(10) and the C(7) hydroxyl groups.

SUMMARY OF THE INVENTION

Among the objects of the present invention, therefore, is the provision of highly efficient processes for the preparation of taxol and other taxanes through selective derivatization of the C(7) group or the C(10) hydroxyl group of 10-DAB and other taxanes, particularly a process in which the C(10) hydroxyl group is protected

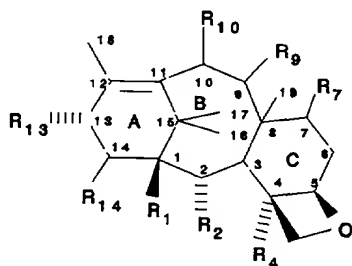
or derivatized prior to the C(7) hydroxyl group; and the provision of C(7) or C(10) derivatized taxanes.

Briefly, therefore, the present invention is directed to a process for the acylation of the C(10) hydroxy group of a taxane. The process comprises forming a reaction mixture containing the taxane and an acylating agent which contains less than one equivalent of an amine base for each equivalent of taxane, and allowing the taxane to react with the acylating agent to form a C(10) acylated taxane.

The present invention is further directed to a process for the silylation of the C(10) hydroxy group of a taxane having a C(10) hydroxy group. The process comprises treating the taxane with a silylamide or a bissilylamide to form a C(10) silylated taxane.

The present invention is further directed to a process for converting the C(7) hydroxy group of a 10-acyloxy-7-hydroxytaxane to an acetal or ketal. The process comprises treating the 10-acyloxy-7-hydroxytaxane with a ketalizing agent in the presence of an acid catalyst to form a C(10) ketalized taxane.

The present invention is further directed to a taxane having the structure:



wherein

M is a metal or comprises ammonium:

R₁ is hydrogen, hydroxy, protected hydroxy, or together with R₁₄ or R₂ forms a carbonate;

R_2 is keto, $-OT_2$, acyloxy, or together with R_1 forms a carbonate;

R_4 is $-OT_4$ or acyloxy;

R_7 is $-OSiR_JR_KR_L$;

R_9 is hydrogen, keto, $-OT_9$, or acyloxy;

R_{10} is hydrogen, keto, $-OT_{10}$, or acyloxy;

R_{13} is hydroxy, protected hydroxy, keto, or $MO-$;

R_{14} is hydrogen, $-OT_{14}$, acyloxy, or together with R_1 forms a carbonate;

R_J , R_K , R_L are independently hydrocarbyl, substituted hydrocarbyl, or heteroaryl, provided, however, if each of R_J , R_K and R_L are alkyl, at least one of R_J , R_K and R_L comprises a carbon skeleton having at least four carbon atoms; and

T_2 , T_4 , T_9 , T_{10} , and T_{14} are independently hydrogen or hydroxy protecting group.

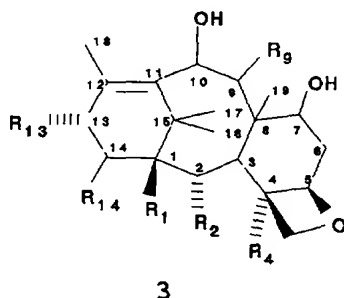
Other objects and features of this invention will be in part apparent and in part pointed out hereinafter.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Among other things, the present invention enables the selective derivatization of the C(10) hydroxyl group of a taxane without first protecting the C(7) hydroxyl group. Stated another way, it has been discovered that the reactivities previously reported for the C(7) and C(10) hydroxyl groups can be reversed, that is, the reactivity of the C(10) hydroxyl group becomes greater than the reactivity of the C(7) hydroxyl group under certain conditions.

Although the present invention may be used to selectively derivatize a taxane having a hydroxy group at C(7) or C(10), it offers particular advantages in the selective derivatization of taxanes having hydroxy groups at C(7) and C(10), i.e., 7,10-dihydroxy taxanes. In

general, 7,10-dihydroxytaxanes which may be selectively derivatized in accordance with the present invention correspond to the following structure:



wherein

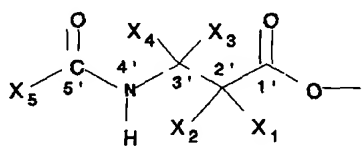
R_1 is hydrogen, hydroxy, protected hydroxy, or together with R_{14} or R_2 forms a carbonate;

R_2 is keto, $-OT_2$, acyloxy, or together with R_1 forms a carbonate;

R_4 is $-OT_4$ or acyloxy;

R_9 is hydrogen, keto, $-OT_9$, or acyloxy;

R_{13} is hydroxy, protected hydroxy, keto, or



R_{14} is hydrogen, $-OT_{14}$, acyloxy or together with R_1 forms a carbonate;

T_2 , T_4 , T_9 , and T_{14} are independently hydrogen or hydroxy protecting group;

X_1 is $-OX_6$, $-SX_7$, or $-NX_8X_9$;

X_2 is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

X_3 and X_4 are independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

X_5 is $-X_{10}$, $-OX_{10}$, $-SX_{10}$, $-NX_8X_{10}$, or $-SO_2X_{11}$;

X_6 is hydrocarbyl, substituted hydrocarbyl, heteroaryl, hydroxy protecting group or a functional group which increases the water solubility of the taxane derivative;

X_7 is hydrocarbyl, substituted hydrocarbyl, heteroaryl, or sulphydryl protecting group;

X_8 is hydrogen, hydrocarbyl, or substituted hydrocarbyl;

X_9 is an amino protecting group;

X_{10} is hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

X_{11} is hydrocarbyl, substituted hydrocarbyl, heteroaryl, $-OX_{10}$, or $-NX_8X_{14}$; and

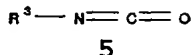
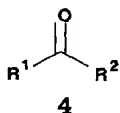
X_{14} is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

Selective C(10) Derivatization

In accordance with the process of the present invention, it has been discovered that the C(10) hydroxyl group of a taxane can be selectively acylated in the absence of an amine base. Preferably, therefore, amine bases such as pyridine, triethylamine, dimethylaminopyridine and 2,6-lutidine, if present at all, are present in the reaction mixture in relatively low concentration. Stated another way, if an amine base is present in the reaction mixture, the molar ratio of the amine base to the taxane is preferably less than 1:1, more preferably less than 10:1, and most preferably less than 100:1.

Acylating agents which may be used for the selective acylation of the C(10) hydroxyl group of a taxane include anhydrides, dicarbonates, thiodicarbonates, and isocyanates. In general, the anhydrides, dicarbonates,

and thiodicarbonates correspond to structure 4 and the isocyanates correspond to structure 5:



wherein R^1 is $-\text{OR}^a$, $-\text{SR}^a$, or R^a ; R^2 is $-\text{OC}(\text{O})\text{R}^b$, $-\text{OC}(\text{O})\text{OR}^b$, $-\text{OC}(\text{O})\text{SR}^b$, $-\text{OPOR}^b\text{R}^c$, or $-\text{OS}(\text{O})_2\text{R}^b$; R^3 is hydrocarbyl, substituted hydrocarbyl, or heteroaryl; and R^a , R^b , R^c are independently hydrocarbyl, substituted hydrocarbyl, or heteroaryl. For example, suitable carboxylic acid anhydride acylating agents include acetic anhydride, chloroacetic anhydride, propionic anhydride, benzoic anhydride, and other carboxylic acid anhydrides containing substituted or unsubstituted hydrocarbyl or heteroaryl moieties; suitable dicarbonate acylating reagents include dibenzyl dicarbonate, diallyl dicarbonate, dipropyl dicarbonate, and other dicarbonates containing substituted or unsubstituted hydrocarbyl or heteroaryl moieties; and suitable isocyanate acylating agents include phenyl isocyanate, and other isocyanates containing substituted or unsubstituted hydrocarbyl or heteroaryl moieties. In addition, although the anhydrides, dicarbonates, and thiodicarbonates used as acylating agents may be mixed, it is generally preferred that they be symmetrical; that is, R^1 and R^2 are selected such that the molecule is symmetrical (e.g., if R^1 is R^a , R^2 is $-\text{OC}(\text{O})\text{R}^b$ with R^a being the same as R^b).

While the acylation of the C(10) hydroxy group of the taxane will proceed at an adequate rate for many acylating agents, it has been discovered that the reaction rate may be increased by including a Lewis acid in the reaction mixture. The concentration of the Lewis

acid appears not to be narrowly critical; experimental evidence obtained to date suggests it may be present in either a stoichiometric or a catalytic amount. In general, Lewis acids which may be used include triflates and halides of elements of groups IB, IIB, IIIB, IVB, VB, VIB, VIIB, VIII, IIIA, IVA, lanthanides, and actinides of the Periodic Table (American Chemical Society format). Preferred Lewis acids include zinc chloride, stannic chloride, cerium trichloride, cuprous chloride, lanthanum trichloride, dysprosium trichloride, and ytterbium trichloride. Zinc chloride or cerium trichloride is particularly preferred when the acylating agent is an anhydride or dicarbonate. Cuprous chloride is particularly preferred when the acylating agent is an isocyanate.

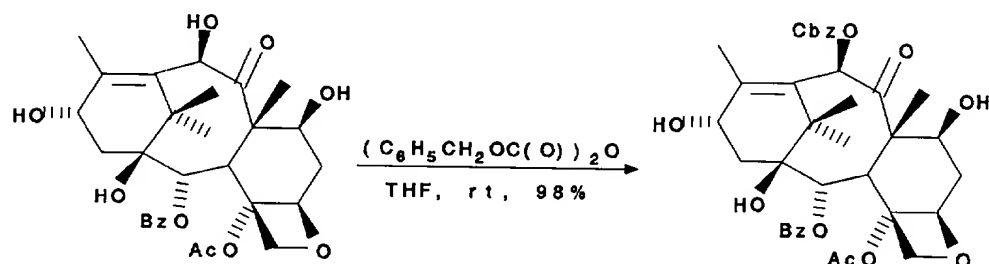
The solvent for the selective acylation is preferably an ethereal solvent such as tetrahydrofuran. Alternatively, however, other solvents such as ether or dimethoxyethane may be used.

The temperature at which the C(10) selective acylation is carried out is not narrowly critical. In general, however, it is preferably carried out at room temperature or higher in order for the reaction to proceed at a sufficiently high rate.

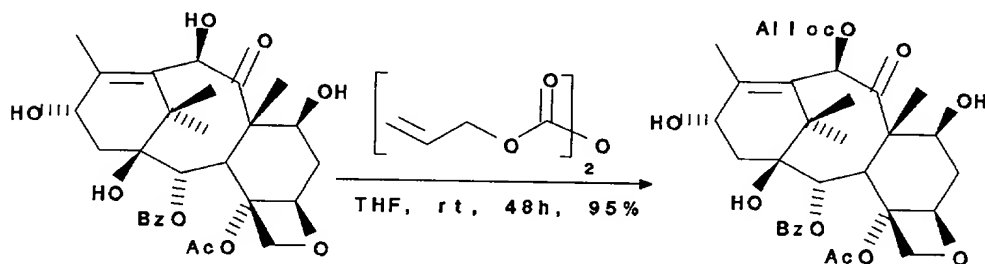
For purposes of illustration, acylating reactions involving dibenzyl dicarbonate, diallyl dicarbonate, acetic anhydride, chloroacetic anhydride and phenyl isocyanate are illustrated in Reaction Schemes 1 through 5 below. In this series of reaction schemes, the taxane which is selectively acylated at the C(10) position is 10-deacetylbaccatin III. It should be understood, however, that these reaction schemes are merely illustrative and that other taxanes having a C(10) hydroxy group, in general, and other

7,10-dihydroxytaxanes, in particular, may be selectively acylated with these and other acylating agents in accordance with the present invention.

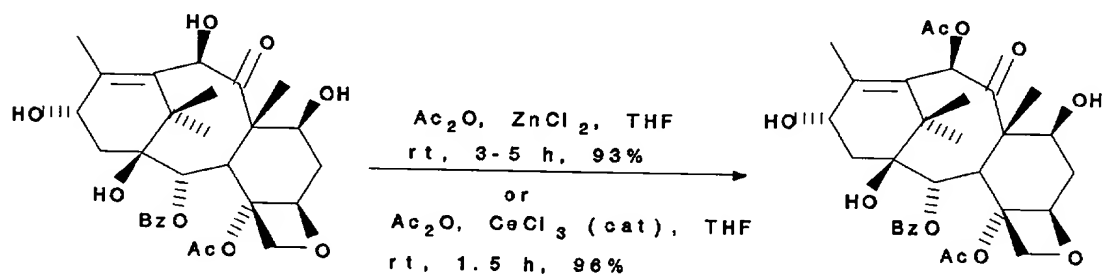
Scheme 1



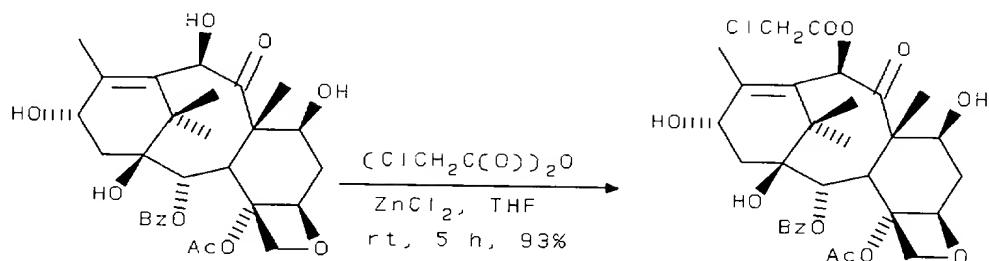
Scheme 2



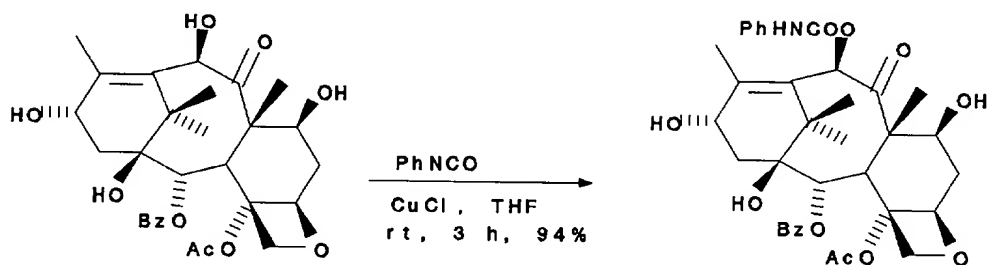
Scheme 3



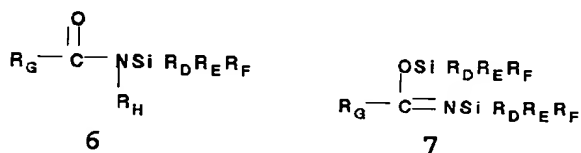
Scheme 4



Scheme 5



In another aspect of the present invention, the C(10) hydroxyl group of a taxane may be selectively silylated. In general, the silylating agent is selected from the group consisting of silylamides and bissilylamides. Preferred silylamides and bissilylamides correspond to structures 6 and 7, respectively:



wherein R_D , R_E , R_F , R_G , and R_H are independently hydrocarbyl, substituted hydrocarbyl, or heteroaryl. Preferably, the silylating agents are selected from the group consisting of tri(hydrocarbyl)silyl-

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trifluoromethylacetamides and bis tri(hydrocarbyl)-silyltrifluoromethylacetamides, with the hydrocarbyl moiety being substituted or unsubstituted alkyl or aryl. For example, the preferred silylamides and bissilylamides include N,O-bis-(trimethylsilyl) trifluoroacetamide, N,O-bis-(triethylsilyl)trifluoroacetamide, N-methyl-N-triethylsilyltrifluoroacetamide, and N,O-bis(t-butyl dimethylsilyl)trifluoroacetamide.

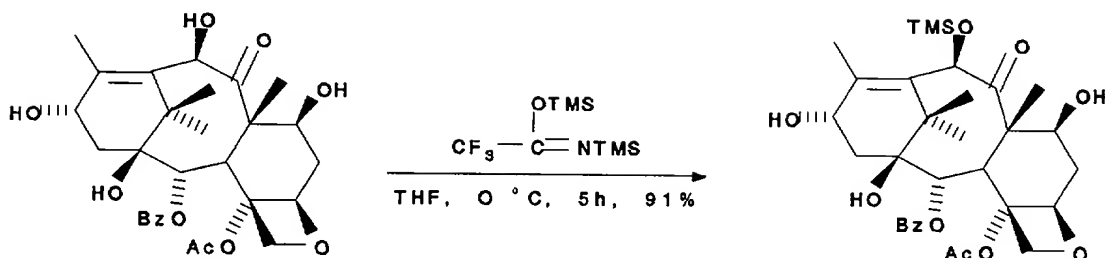
The silylating agents may be used either alone or in combination with a catalytic amount of a base such as an alkali metal base. Alkali metal amides, such as lithium amide catalysts, in general, and lithium hexamethyldisilazide, in particular, are preferred.

The solvent for the selective silylation reaction is preferably an ethereal solvent such as tetrahydrofuran. Alternatively, however, other solvents such as ether or dimethoxyethane may be used.

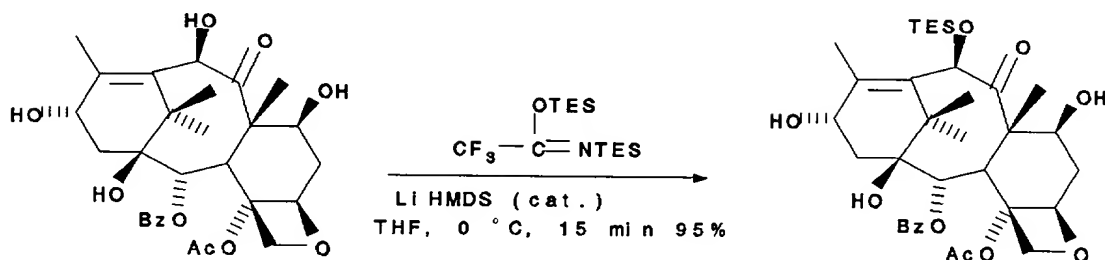
The temperature at which the C(10) selective silylation is carried out is not narrowly critical. In general, however, it is carried out at 0 °C or greater.

Selective C(10) silylation reactions involving N,O-bis(trimethylsilyl)trifluoroacetamide and N,O-bis(triethylsilyl)trifluoroacetamide are illustrated in Reaction Schemes 6 and 7 below. In these reaction schemes, the taxane which is selectively silylated at the C(10) position is 10-deacetylbaccatin III. It should be understood, however, that these reaction schemes are merely illustrative and that other taxanes, having a C(10) hydroxy group, in general, and other 7,10-dihydroxytaxanes, in particular, may be selectively silylated with these and other silylating agents in accordance with the present invention.

Scheme 6



Scheme 7



After the C(10) hydroxyl group of a 7,10-dihydroxytaxane has been derivatized as described herein, the C(7) hydroxyl group can readily be protected or otherwise derivatized selectively in the presence of the C(1) and C(13) hydroxyl groups (and a C(14) hydroxy group, if present).

Selective C(7) Derivatization

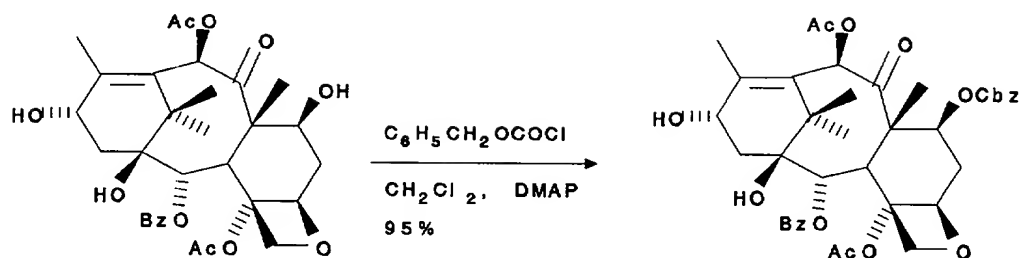
Selective acylation of the C(7) hydroxyl group of a C(10) acylated or silylated taxane can be achieved using any of a variety of common acylating agents including, but not limited to, substituted and unsubstituted carboxylic acid derivatives, e.g., carboxylic acid halides, anhydrides, dicarbonates, isocyanates and haloformates. For example, the C(7) hydroxyl group of

baccatin III, 10-acyl-10-deacetyl baccatin III or 10-trihydrocarbylsilyl-10-deacetyl baccatin III can be selectively acylated with dibenzyl dicarbonate, diallyl dicarbonate, 2,2,2-trichloroethyl chloroformate, benzyl chloroformate or another common acylating agent.

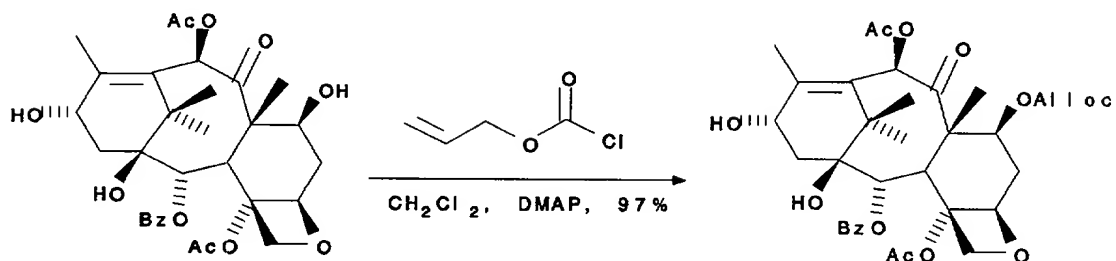
In general, acylation of the C(7) hydroxy group of a C(10) acylated or silylated taxane are more efficient and more selective than are C(7) acylations of a 7,10-dihydroxy taxane such as 10-DAB, i.e., once the C(10) hydroxyl group has been acylated or silylated, there is a significant difference in the reactivity of the remaining C(7), C(13), and C(1) hydroxyl groups (and the C(14) hydroxyl group, if present). These acylation reactions may optionally be carried out in the presence or absence of an amine base.

Examples of selective C(7) acylation of a taxane having an acylated or silylated C(10) hydroxy group are shown in Reaction Schemes 8 through 11. In these reaction schemes, the taxane which is selectively acylated at the C(7) position is baccatin III or 10-triethylsilyl-10-deacetyl baccatin III. It should be understood, however, that these reaction schemes are merely illustrative and that taxanes having other acyl and silyl moieties at C(10) as well as other substituents at other taxane ring positions may be selectively acylated at C(7) with these and other acylating agents in accordance with the present invention.

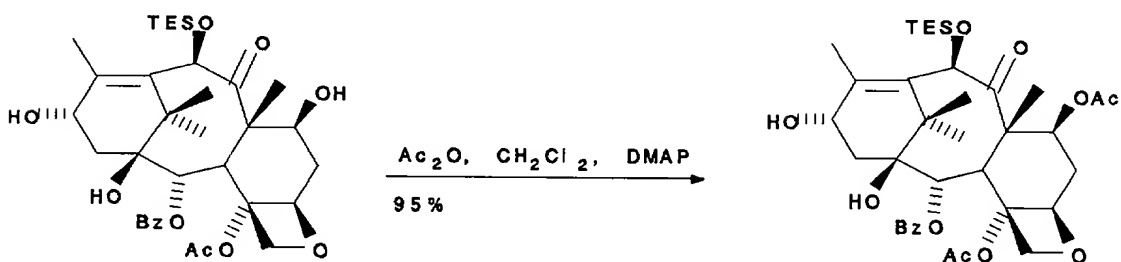
Scheme 8



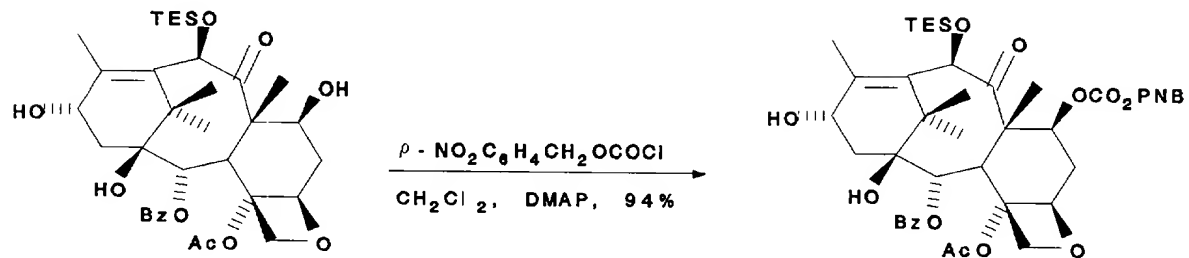
Scheme 9



Scheme 10



Scheme 11



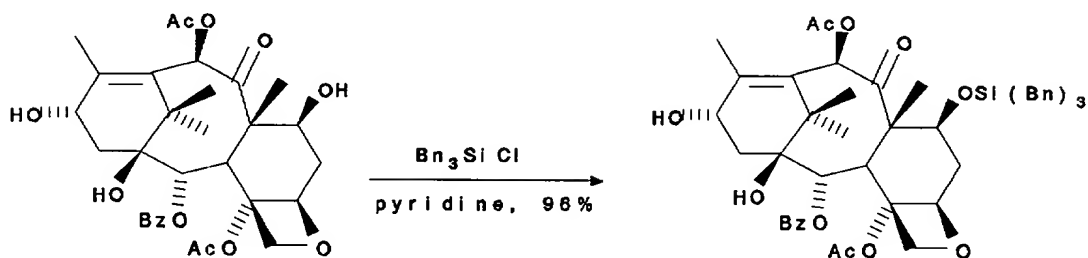
Alternatively, the C(7) hydroxyl group of a C(10) acylated taxane derivative can be selectively protected using any of a variety of hydroxy protecting groups, such as acetal, ketal, silyl, and removable acyl protecting groups. For example, the C(7) hydroxyl group may be silylated using any of a variety of common silylating agents including, but not limited to, tri(hydrocarbyl)silyl halides and tri(hydrocarbyl)silyl triflates. The hydrocarbyl moieties of these compounds may be substituted or unsubstituted and preferably are substituted or unsubstituted alkyl or aryl. For example, the C(7) hydroxyl group of baccatin III can be selectively silylated using silylating agents such as tribenzylsilyl chloride, trimethylsilyl chloride, triethylsilyl chloride, dimethyl isopropylsilyl chloride, dimethyl phenylsilyl chloride, and the like.

In general, silylations of the C(7) hydroxy group of a C(10) acylated taxanes are more efficient and more selective than are silylations of a 7,10-dihydroxy taxane such as 10-DAB, i.e., once the C(10) hydroxyl group has been acylated, there is a significant difference in the reactivity of the remaining C(7), C(13), and C(1) hydroxyl groups (and the C(14) hydroxyl group, if present). The C(7) silylation reaction may be carried out under a wide range of conditions, including in the presence or absence of an amine base.

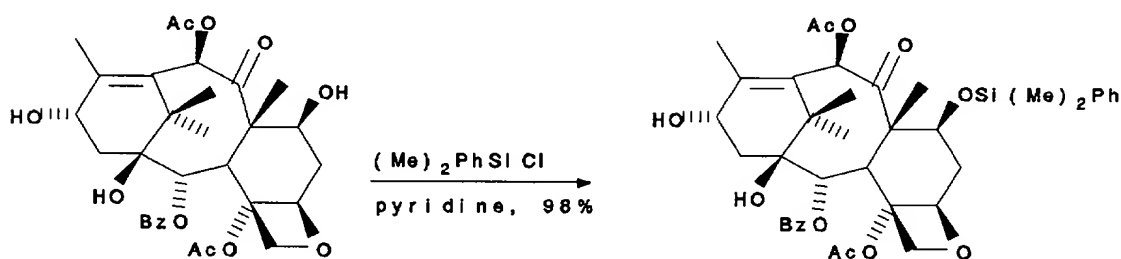
Examples of selective C(7) silylation of C(10) acylated taxanes are shown in Reaction Schemes 12 through 15. In these reaction schemes, the taxane which is selectively silylated at the C(7) position is baccatin III or another C(10)-acyloxy derivative of 10-deacetyl baccatin III. It should be understood, however, that these reaction schemes are merely illustrative and that other taxanes may be selectively silylated with

these and other silylating agents in accordance with the present invention.

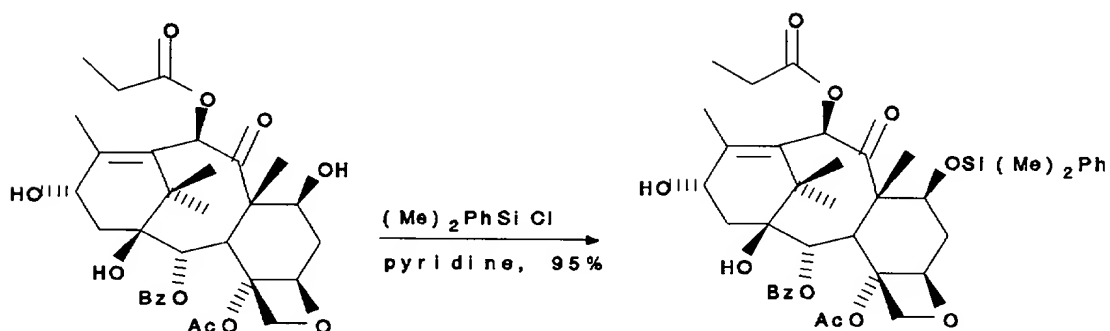
Scheme 12



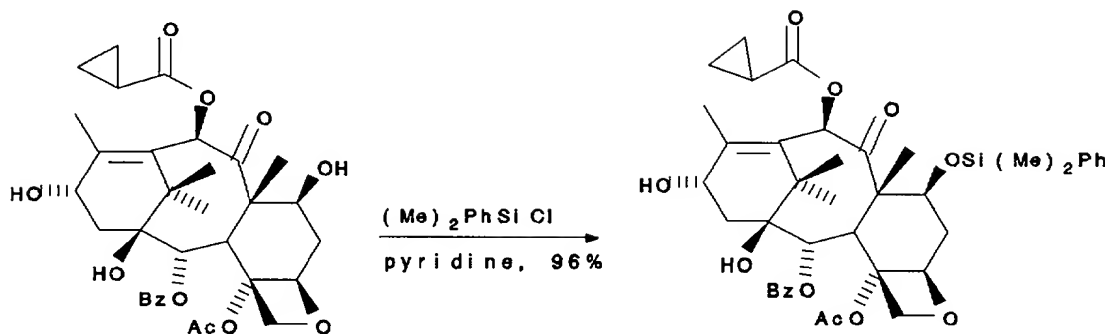
Scheme 13



Scheme 14



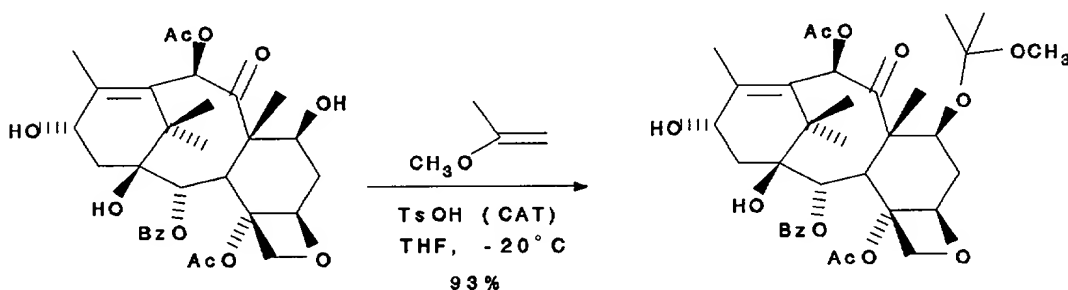
Scheme 15



Alternatively, the C(7) hydroxyl group of C(10) acylated taxanes can be selectively protected using any of a variety of common reagents including, but not limited to, simple acetals, ketals and vinyl ethers, in the presence of an acid catalyst. These reagents (whether acetal, ketal, vinyl ether or otherwise) are referred to herein as "ketalizing agents" and are described in "Protective Groups in Organic Synthesis" by T. W. Greene, John Wiley and Sons, 1981. The acid catalyst used may be an organic or inorganic acid, such as toluenesulfonic acid or camphorsulfonic acid, in at least a catalytic amount. For example, the C(7) hydroxyl group of baccatin III can be selectively ketalized using 2-methoxy propene. Other suitable reagents for the preparation of acetals and ketals include methyl vinyl ether, ethyl vinyl ether, tetrahydropyran, and the like.

Selective ketalization of the C(7) substituent of a C(10) acylated taxane is more efficient and more selective than it is with 10-DAB, i.e., once the C(10) hydroxyl group has been acylated, there is a large difference in the reactivity of the remaining C(7), C(13), and C(1) hydroxyl groups (and the C(14) hydroxyl group, if present).

Scheme 16

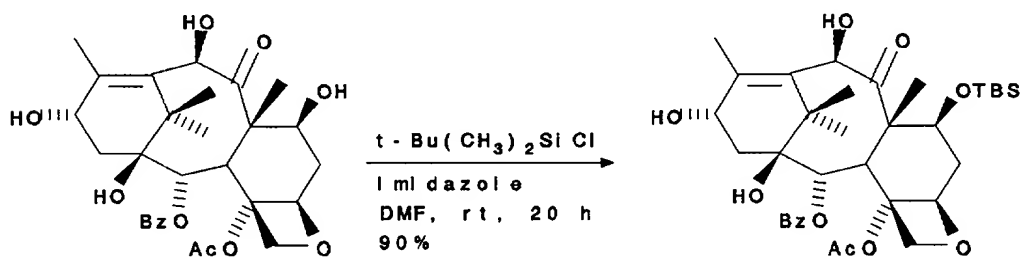


In general, the C(7) hydroxyl group of a taxane can be selectively silylated with a silylating agent which includes the $-SiR_JR_KR_L$ moiety wherein R_J , R_K and R_L are independently substituted or unsubstituted hydrocarbyl or heteroaryl, provided that any substituents are other than hydroxyl. In one embodiment of the present invention, if each of R_J , R_K and R_L is alkyl, then at least one of R_J , R_K , and R_L comprises a carbon skeleton (i.e., carbon chain or ring(s)) having at least four carbon atoms. Suitable silylating agents include silyl halides and silyl triflates, for example, tri(hydrocarbyl)silyl halides and tri(hydrocarbyl)silyl triflates. The hydrocarbyl

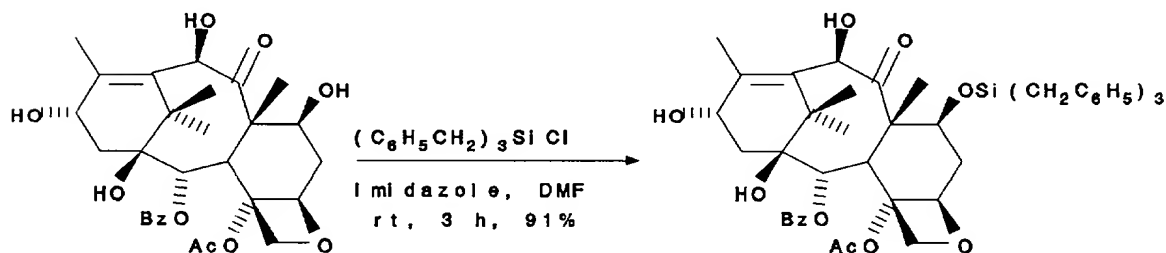
substituents of these silylating agents may be substituted or unsubstituted and preferably are substituted or unsubstituted alkyl or aryl.

The selective silylation of the C(7) hydroxy group may be carried out in a solvent, such as dimethyl formamide ("DMF") or pyridine and in the presence of an amine base, such as imidazole or pyridine. Reaction Schemes 17 - 20 illustrate the silylation of the C(7) hydroxy group of 10-DAB in high yield by treating 10-DAB with t-butyldimethylsilyl chloride, tribenzylsilyl chloride, dimethyl-isopropylsilyl chloride, and dimethylphenylsilyl chloride, respectively. Silylation under these conditions was surprising in view of the report by Denis, et. al. (*J. Am. Chem. Soc.*, **1988**, 110, 5917) that selective formation of 7-TBS-10-DAB was not possible.

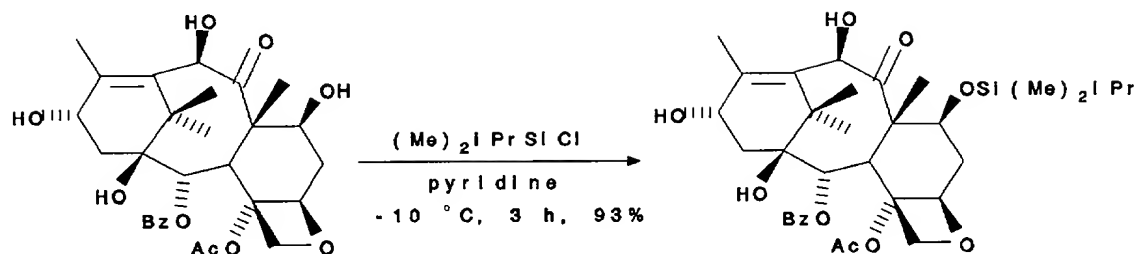
Scheme 17



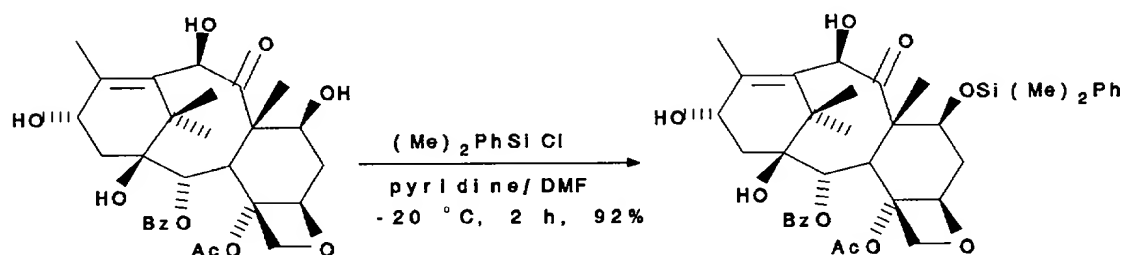
Scheme 18



Scheme 19



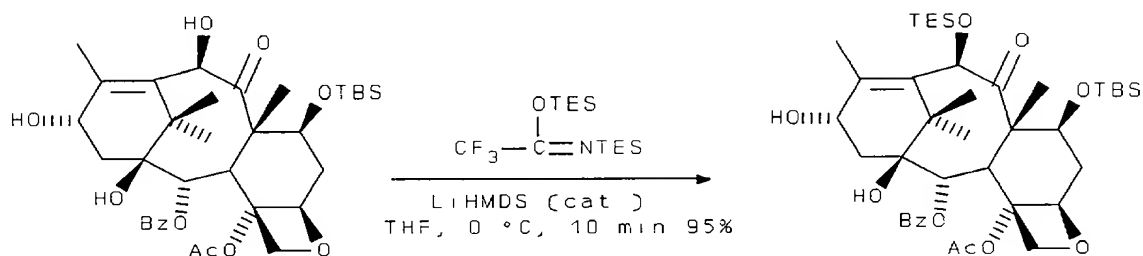
Scheme 20



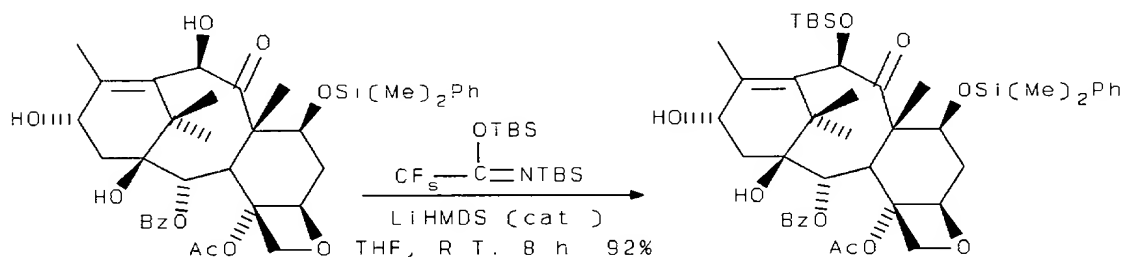
The process of the present invention can also be used to protect the C(7) and C(10) hydroxy groups of a 7,10-dihydroxytaxane with different silyl protecting groups. By selecting groups which can be removed under different conditions, the C(7) and C(10) hydroxy groups can be separately accessed for derivatization. These reactions, therefore, increase the flexibility of the overall process and, enable a higher yield for many of the individual protecting reactions relative to the yield obtained using currently available processes. For example, the triethylsilyl protecting group is more readily removed from C(10) than is the t-butyldimethylsilyl protecting group from C(7) and the dimethylphenylsilyl protecting group is more readily removed from C(7) than is the t-butyldimethylsilyl protecting group from C(10). The preparation of 7-t-

butyldimethylsilyl-10-triethylsilyl-10-DAB and 7-dimethylphenylsilyl-10-t-butyldimethylsilyl-10-DAB are illustrated in Reaction Schemes 21 and 22.

Scheme 21



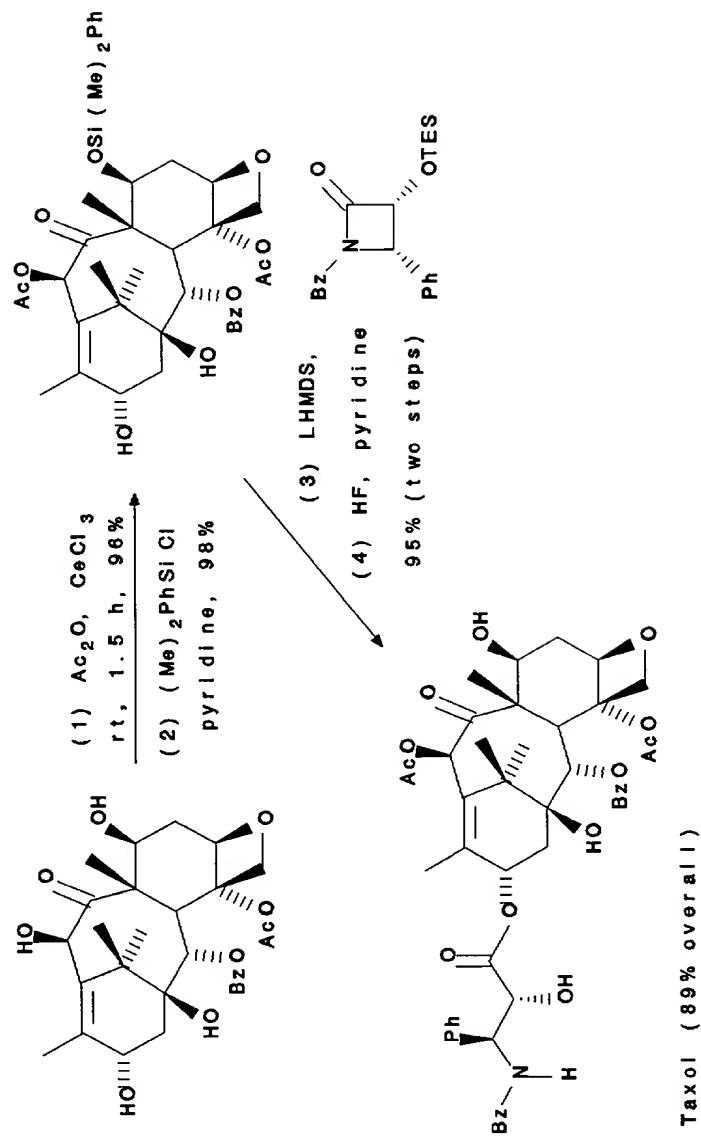
Scheme 22



The methods disclosed herein may be used in connection with a large number of different taxanes obtained from natural or synthetic sources to prepare a wide variety of taxane intermediates which may then be further derivatized. For example, the methods of the present invention may be effectively used to protect the C(7) and/or C(10) hydroxy functional group prior to the coupling reaction between a C(13) side chain precursor and a taxane to introduce a C(13) β -amido ester side chain, and also prior to the reactions for preparing taxanes having alternative substituents at various locations on the taxane nucleus.

The attachment of a C(13) side chain precursor to a taxane may be carried out by various known techniques. For example, a side chain precursor such as an appropriately substituted β -lactam, oxazoline, oxazolidine carboxylic acid, oxazolidine carboxylic acid anhydride, or isoserine derivative may be reacted with a tricyclic or tetracyclic taxane having a C(13) hydroxy, metallic oxide or ammonium oxide substituent to form compounds having a β -amido ester substituent at C(13) as described, for example, Taxol: Science and Applications, M. Suffness, editor, CRC Press (Boca Rotan, FL) 1995, Chapter V, pages 97-121. For example, the synthesis of taxol from 10-DAB is illustrated in reaction scheme 23. It should be noted that while a β -lactam and 10-DAB are used in this reaction scheme, other side chain precursors and other taxanes could be substituted therefor without departing from the present invention.

Scheme 23

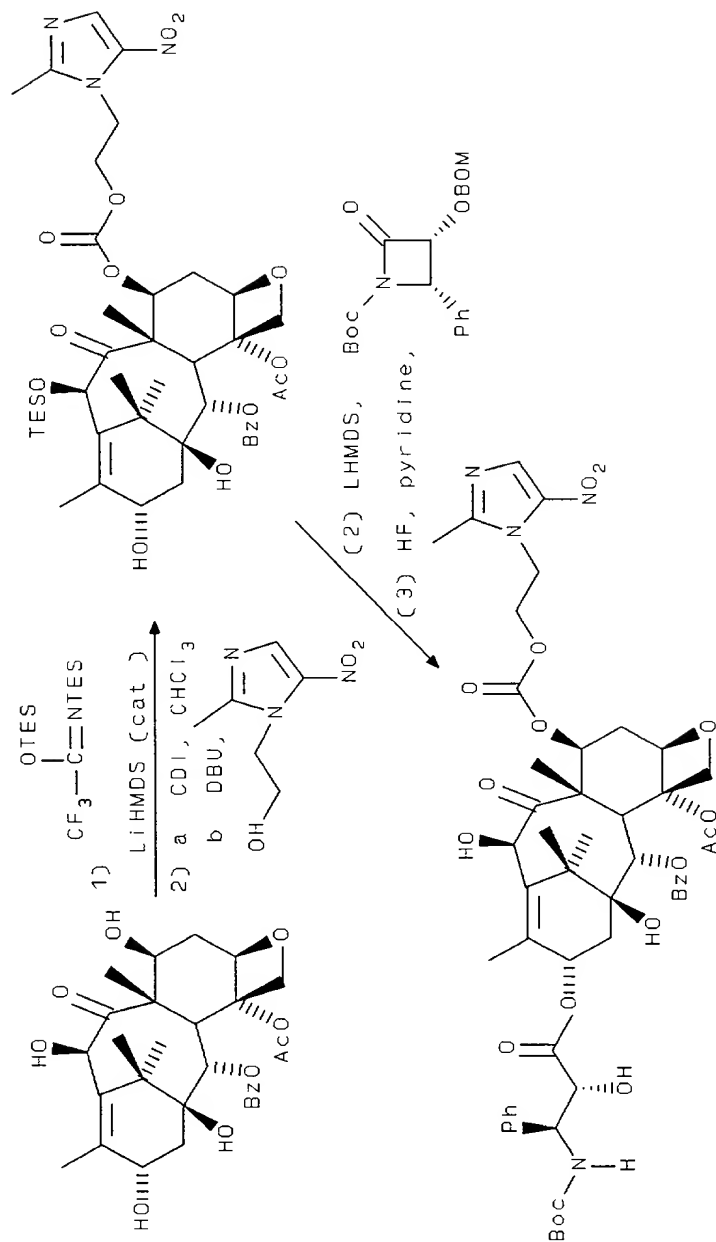


The process illustrated in Reaction Scheme 23 is significantly more efficient than any other currently known process, due to the high yields and selectivity of the cerium trichloride catalyzed acetylation of the C(10) hydroxyl group of 10-DAB and the subsequent silylation of the C(7) hydroxyl group. The synthesis proceeds in four steps and 89% overall yield.

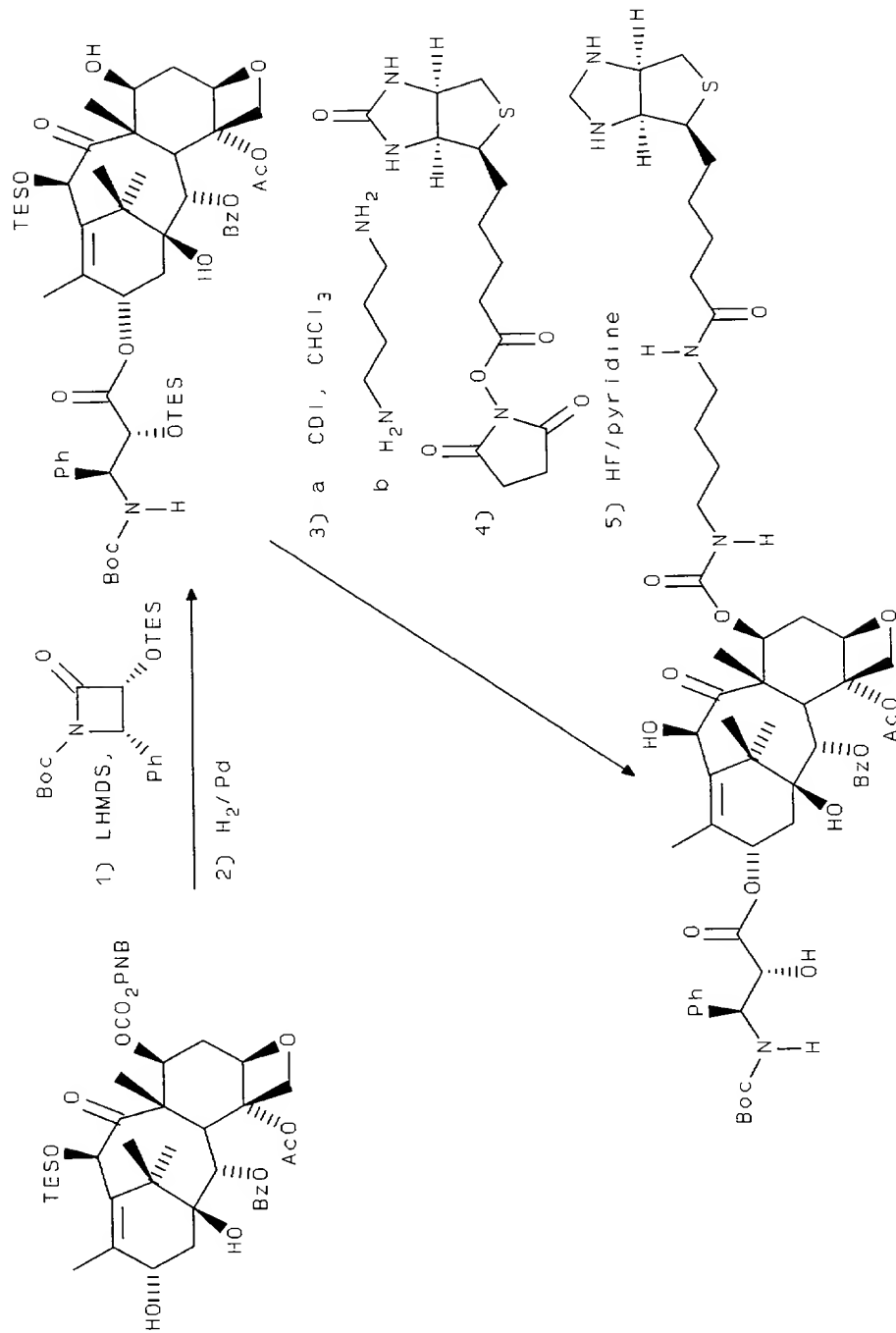
Reaction schemes 24 and 25 illustrate the preparation of taxanes having substituents appended to the C(7) hydroxyl group and a free C(10) hydroxyl group. The method of the current invention provides flexibility so that the substituent attached to the C(7) hydroxyl group can be put in place either before or after attachment of the C(13) side chain.

Reaction scheme 24 outlines the preparation of a taxane which has been found to be a potent chemotherapeutic radiosensitizer, illustrating attachment of the substituent at the C(7) hydroxyl group before introduction of the C(13) side chain. According to the process of reaction scheme 7, 10-DAB is first converted to 10-TES-10-DAB. The C(7) hydroxyl group is then converted to an intermediate imidazolide by treatment with carbonyl diimidazole, and the intermediate imidazolide subsequently reacts, without isolation, with metronidazole alcohol to provide 7-metro-10-TES-10-DAB. Coupling of 7-metro-10-TES-10-DAB with a β -lactam to introduce the side chain at C(13) is followed by removal of the TES groups at C(10) and C(2') by treatment with HF and pyridine.

Scheme 24



Scheme 25



The protected taxane derivatives or the intermediates or starting materials used in the preparation of such protected taxane derivatives can be further modified to provide for alternative substituents at various positions of the taxane.

Taxanes having C(2) and/or C(4) substituents other than benzoyloxy and acetoxy, respectively, can be prepared from baccatin III, 10-DAB and other taxanes as more fully described in PCT Patent Application WO 94/01223. In general, the C(2) and C(4) acyloxy substituents are treated with lithium aluminum hydride or another suitable reducing agent to form hydroxy groups at C(2) and C(4) which may then be reacted, for example, with carboxylic acid halides (optionally after protection of the C(2) hydroxy group together with the C(1) hydroxy group with a 1,2-carbonate protecting group) to obtain the desired C(2) and C(4) derivatives.

Taxanes having C(7) substituents other than hydroxy and acyloxy as described herein can be prepared from baccatin III, 10-DAB, and other taxanes as more fully described in PCT Patent Application WO 94/17050. For example, a C(7) xanthate may be subjected to tin hydride reduction to yield the corresponding C(7) dihydro taxane. Alternatively, C(7) fluoro-substituted taxanes can be prepared by treatment of C(13)-triethylsilyl-protected baccatin III with 2-chloro-1,1,2-trifluorotriethylamine at room temperature in THF solution. Other baccatin derivatives with a free C(7) hydroxyl group behave similarly. Alternatively, 7-chloro baccatin III can be prepared by treatment of baccatin III with methanesulfonyl chloride and triethylamine in methylene chloride solution containing an excess of triethylamine hydrochloride.

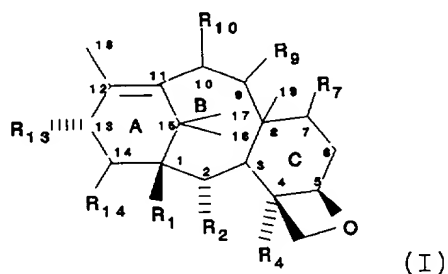
Taxanes having C(9) substituents other than keto can be prepared from baccatin III, 10-DAB and other taxanes as more fully described in PCT Patent Application WO 94/20088. In general, the C(9) keto substituent of the taxane is selectively reduced to yield the corresponding C(9) β -hydroxy derivative with a borohydride, preferably tetrabutylammonium borohydride (Bu_4NBH_4) or triacetoxy-borohydride. The C(9) β -hydroxy derivative can then be protected at C(7) with a hydroxy protecting group and the C(9) hydroxy group can be acylated following the methods described herein for acylation of the C(7) hydroxy group. Alternatively, reaction of 7-protected-9 β -hydroxy derivative with KH causes the acetate group (or other acyloxy group) to migrate from C(10) to C(9) and the hydroxy group to migrate from C(9) to C(10), thereby yielding a 10-desacetyl derivative, which can be acylated as described elsewhere herein.

Taxanes having C(10) substituents other than hydroxy, acyloxy or protected hydroxy as described herein may be prepared as more fully described in PCT Patent Application WO 94/15599 and other literature references. For example, taxanes having a C(10) keto substituent can be prepared by oxidation of 10-desacetyl taxanes. Taxanes which are dihydro substituted at C(10) can be prepared by reacting a C(10) hydroxy or acyloxy substituted taxane with samarium diiodide.

Taxanes having a C(14) substituent other than hydrogen may also be prepared. The starting material for these compounds may be, for example, a hydroxylated taxane (14-hydroxy-10-deacetyl baccatin III) which has been discovered in an extract of yew needles (C&EN, p 36-37, April 12, 1993). Derivatives of this hydroxylated taxane having the various C(2), C(4), C(7), C(9), C(10), C3' and C5' functional groups described above may also be

prepared by using this hydroxylated taxane. In addition, the C(14) hydroxy group together with the C(1) hydroxy group of 10-DAB can be converted to a 1,2-carbonate as described in C&EN or it may be converted to a variety of esters or other functional groups as otherwise described herein in connection with the C(2), C(4), C(9) and C(10) substituents.

The process of the present invention thus enables the preparation of taxanes having the following structure:



wherein

M comprises ammonium or is a metal;

R₁ is hydrogen, hydroxy, protected hydroxy, or together with R₁₄ or R₂ forms a carbonate;

R₂ is keto, -OT₂, acyloxy, or together with R₁ forms a carbonate;

R₄ is -OT₄ or acyloxy;

R₇ is hydrogen, halogen, -OT₇, or acyloxy;

R₉ is hydrogen, keto, -OT₉, or acyloxy;

R₁₀ is hydrogen, keto, -OT₁₀, or acyloxy;

R₇, R₉, and R₁₀ independently have the alpha or beta stereochemical configuration;

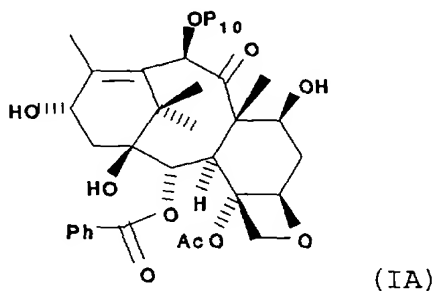
R₁₃ is hydroxy, protected hydroxy, keto, MO- or



In one embodiment of the present invention, the substituents of the taxane (other than the C(7), C(10) and C(13) substituents) correspond to the substituents present on baccatin III or 10-DAB. That is, R₁₄ is

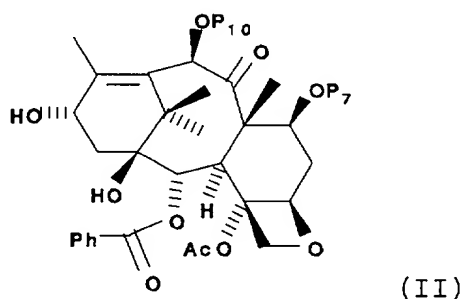
hydrogen, R_9 is keto, R_4 is acetoxy, R_2 is benzyloxy, and R_1 is hydroxy. In other embodiments, the taxane has a structure which differs from that of taxol or Taxotere® with respect to the C(13) side chain and at least one other substituent. For example, R_{14} may be hydroxy; R_2 may be hydroxy, $-\text{OCOZ}_2$ or $-\text{OCOOZ}_{22}$ wherein Z_2 is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl and Z_{22} is hydrocarbyl, substituted hydrocarbyl, or heteroaryl; R_4 may be hydroxy, $-\text{OCOZ}_4$ or $-\text{OCOOZ}_{44}$ wherein Z_4 is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl and Z_{44} is hydrocarbyl, substituted hydrocarbyl, or heteroaryl; R_7 may be hydrogen, hydroxy, $-\text{OCOZ}_7$ or $-\text{OCOOZ}_{77}$ wherein Z_7 is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl and Z_{77} is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl, R_9 may be hydrogen, hydroxy, $-\text{OCOZ}_9$ or $-\text{OCOOZ}_{99}$ wherein Z_9 is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl and Z_{99} is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl, and R_{10} may be hydrogen, hydroxy, $-\text{OCOZ}_{10}$ or $-\text{OCOOZ}_{1010}$ wherein Z_{10} is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl and Z_{1010} is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

In a preferred embodiment, the taxane has the formula



wherein P_{10} is acyl, said acyl comprising at least three carbon atoms or two carbon atoms and a nitrogen, oxygen or sulfur atom. Stated another way, $-OP_{10}$ is other than acetoxy. More preferably, P_{10} is $-(C=O)R_A$, $-(C=O)OR_B$, or $-(C=O)NR_C$ wherein R_A is substituted or unsubstituted hydrocarbyl or heteroaryl, said unsubstituted hydrocarbyl comprising at least two carbon atoms; R_B and R_C are independently substituted or unsubstituted hydrocarbyl. Still more preferably, R_A is substituted or unsubstituted alkyl or aryl, said unsubstituted alkyl comprising at least two carbon atoms; and R_B and R_C are independently substituted or unsubstituted alkyl or aryl.

In another embodiment of the invention, the taxane has the formula



wherein P_7 and P_{10} are independently substituted or unsubstituted acyl. In this embodiment, P_7 and P_{10} are preferably different.

Definitions

As used herein, the terms "selective" and "selective derivatization" shall mean that the desired product is preferentially formed over any other by-product. Preferably, the desired product is present in a molar ratio of at least 9:1 relative to any other by-product and, more preferably, is present in a molar ratio of at least 20:1 relative to any other by-product.

In addition, "Ph" means phenyl; "Bz" means benzoyl; "Bn" means benzyl; "Me" means methyl; "Et" means ethyl; "iPr" means isopropyl; "tBu" and "t-Bu" means tert-butyl; "Ac" means acetyl; "TES" means triethylsilyl; "TMS" means trimethylsilyl; "TBS" means Me₂t-BuSi-; "CDI" means carbonyl diimidazole; "BOM" means benzyloxymethyl; "DBU" means diazabicycloundecane; "DMAP" means p-dimethylamino pyridine; "LHMDS" or "LiHMDS" means lithium hexamethyldisilazide; "DMF" means dimethylformamide; "10-DAB" means 10-desacetylbaecatin III; "Cbz" means benzyloxycarbonyl; "Alloc" means allyloxycarbonyl; "THF" means tetrahydrofuran; "BOC" means benzyloxycarbonyl; "PNB" means para-nitrobenzyl; "Troc" means 2,2,2-trichloroethoxycarbonyl; "EtOAc" means ethyl acetate; "THF" means tetrahydrofuran; "protected hydroxyl" means -OP wherein P is a hydroxyl protecting group; and "hydroxyl protecting group" includes, but is not limited to, acetals having two to ten carbons, ketals having two to ten carbons, and ethers, such as methyl, t-butyl, benzyl, p-methoxybenzyl, p-nitrobenzyl, allyl, trityl, methoxymethyl, methoxyethoxymethyl, ethoxyethyl, methoxypropyl, tetrahydropyranyl, tetrahydrothiopyranyl; and trialkylsilyl ethers such as trimethylsilyl ether, triethylsilyl ether, dimethylarylsilyl ether, triisopropylsilyl ether and t-butyltrimethylsilyl ether; esters such as benzoyl, acetyl, phenylacetyl, formyl, mono-, di-, and trihaloacetyl such as chloroacetyl, dichloroacetyl, trichloroacetyl, trifluoroacetyl; and carbonates including but not limited to alkyl carbonates having from one to six carbon atoms such as methyl, ethyl, n-propyl, isopropyl, n-butyl, t-butyl; isobutyl, and n-pentyl; alkyl carbonates having from one to six carbon atoms and substituted with one or more halogen atoms such as 2,2,2-trichloroethoxymethyl and

[illegible]

The alkyl groups described herein are preferably lower alkyl containing from one to six carbon atoms in the principal chain and up to 20 carbon atoms. They may be straight, branched chain or cyclic and include methyl, ethyl, propyl, isopropyl, butyl, hexyl and the like. They may be substituted with aliphatic or cyclic hydrocarbyl radicals.

The alkenyl groups described herein are preferably lower alkenyl containing from two to six carbon atoms in the principal chain and up to 20 carbon atoms. They may be straight or branched chain and include ethenyl, propenyl, isopropenyl, butenyl, isobutenyl, hexenyl, and the like. They may be substituted with aliphatic or cyclic hydrocarbyl radicals.

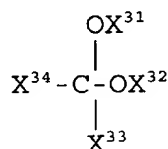
The alkynyl groups described herein are preferably lower alkynyl containing from two to six carbon atoms in

The substituted hydrocarbyl moieties described herein are hydrocarbyl moieties which are substituted with at least one atom other than carbon and hydrogen, including moieties in which a carbon chain atom is substituted with a hetero atom such as nitrogen, oxygen, silicon, phosphorous, boron, sulfur, or a halogen atom. These substituents include hydroxy; lower alkoxy such as methoxy, ethoxy, butoxy; halogen such as chloro or fluoro; ethers; acetals; ketals; esters; heteroaryl such

as furyl or thienyl; alkanoxy; acyl; acyloxy; nitro; amino; and amido.

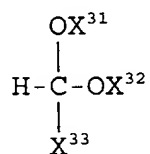
The acyl moieties and the acyloxy moieties described herein contain hydrocarbyl, substituted hydrocarbyl or heteroaryl moieties. In general, they have the formulas $-C(O)G$ and $-OC(O)G$, respectively, wherein G is substituted or unsubstituted hydrocarbyl, hydrocarbyloxy, hydrocarbylamino, hydrocarbylthio or heteroaryl.

The ketal moieties described herein have the general formula



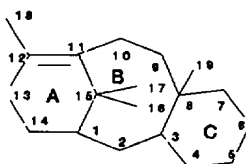
wherein X^{31} , X^{32} , X^{33} and X^{34} are independently hydrocarbyl, substituted hydrocarbyl or heteroaryl moieties. They may be optionally substituted with the various substituents defined herein. The ketal moieties are preferably substituted or unsubstituted alkyl or alkenyl, and more preferably substituted or unsubstituted lower (C_1-C_6) alkyl. These ketal moieties additionally may encompass sugars or substituted sugars and include ketal moieties prepared from sugars or substituted sugars such as glucose and xylose. When a ketal moiety is incorporated into a taxane of the present invention as a C(7) hydroxy protecting group, then either X^{31} or X^{32} represents the taxane moiety.

The acetal moieties described herein have the general formula



wherein X^{31} , X^{32} and X^{33} are independently hydrocarbyl, substituted hydrocarbyl or heteroaryl moieties. They may be optionally substituted with the various substituents defined herein other than hydroxyl. The acetal moieties are preferably substituted or unsubstituted alkyl or alkenyl, and more preferably substituted or unsubstituted lower (C_1 - C_6) alkyl. These acetal moieties additionally may encompass sugars or substituted sugars and include acetal moieties prepared from sugars or substituted sugars such as glucose and xylose. When an acetal moiety is incorporated into a taxane of the present invention as a C(7) hydroxy protecting group, then either X^{31} or X^{32} represents the taxane moiety.

The term "taxane" as used herein, denotes compounds containing the A, B and C rings (with numbering of the ring positions shown herein):



The following examples illustrate the invention.

Example 1

A. Selective acylation of a C(10) hydroxyl group:

10-Cbz-10-DAB. To a solution of 10-DAB (30 mg, 0.055 mmol) in THF (1 mL) at room temperature was added dibenzyl pyrocarbonate (320 mg, 1.1 mmol, 20 equiv) under N_2 . The reaction mixture was stirred at room temperature for 24 h. EtOAc (10 mL) was added, and the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the

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solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:1) as the eluent and dried in vacuo overnight to give 10-cbz-10-DAB as a colorless solid: yield, 37 mg (98%). mp 205-206 °C; $[\alpha]_{\text{D}}^{25}$ -63° (CHCl₃, c = 0.41); ¹H NMR (400 MHz, CDCl₃) δ 1.11(s, 3 H, Me17), 1.13(s, 3 H, Me16), 1.58(s, 1 H, 1-OH), 1.71(s, 3 H, Me19), 1.89(ddd, J = 14.7, 10.9, 2.3 Hz, 1 H, H6b), 2.00(d, J = 5.1 Hz, 1 H, 13-OH), 2.08(d, J = 1.0, 3 H, Me18), 2.28(s, 3 H, 4-Ac), 2.30(m, 2 H, H14a, H14b), 2.43(d, J = 4.1 Hz, 1 H, 7-OH), 2.58(ddd, J = 14.7, 9.6, 6.6 Hz, 1 H, H6a), 3.88(d, J = 6.9 Hz, 1 H, H3), 4.19(d, J = 8.6 Hz, 1 H, H20b), 4.31(d, J = 8.6 Hz, 1 H, H20a), 4.44(ddd, J = 10.9, 6.6, 4.1 Hz, 1 H, H7), 4.89(m, 1 H, H13), 4.98(dd, J = 9.6, 2.3 Hz, 1 H, H5), 5.23(d, J = 12.1, 1 H, CHH'OC(O)), 5.26(d, J = 12.1, 1 H, CHH'OC(O)), 5.65(d, J = 6.9 Hz, 1 H, H2), 6.19(s, 1 H, H10), 7.35-7.44(m, 5 H, PhCH₂O), 7.48(dd, J = 8.1, 7.6 Hz, 2 H, benzoate, m), 7.60(tt, J = 7.6, 1.0 Hz, 1 H, benzoate, p), 8.11(d, J = 8.1, 1.0 Hz, 2 H, benzoate, o) ppm. ¹³C NMR (75 MHz, CDCl₃) δ 9.1(Me(19)), 15.3(Me(18)), 20.7(4-Ac), 22.3, 26.7(Me16, Me17), 35.5(C(6)), 38.6(C(14)), 42.5(C(15)), 46.1(C(3)), 58.7(C(8)), 67.9(C(13)), 70.5(OCH₂Ph), 72.2, 75.0, 76.5(C(7), C(2), C(20)), 79.0, 79.1 (C(1), C(10)), 80.9(C(4)), 84.5(C(5)), 128.6, 128.8, 129.7, 130.3, 131.9, 133.8(OCH₂Ph, benzoate), 135.1(C(11)), 147.5(C(12)), 155.6(OC(O)O), 167.4(benzoate), 171.0(4-Ac), 204.7(C(9))ppm. Anal. Calcd for C₃₇H₄₂O₁₂ · 1/2H₂O: C, 64.62; H, 6.30. Found: C, 64.34; H, 6.31.

10-Alloc-10-DAB. To a solution of 10-DAB (30 mg, 0.055 mmol) in THF (1 mL) at room temperature was added diallyl pyrocarbonate (366 mL, 2.2 mmol, 40 equiv) under

N₂. The reaction mixture was stirred at room temperature for 48 h. TLC analysis indicated the presence of the desired product along with unreacted starting material. EtOAc (20 mL) was added, and the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated. The residue was purified by flash column chromatography using EtOAc: hexanes (1:1) as the eluent and dried *in vacuo* to overnight to give 10-alloc-10-DAB as a colorless solid: yield, 23 mg (67%, 95% at the conversion of 70%). The recovered 10-DAB, 9 mg (30%). 10-alloc-10-DAB: mp 201-203 °C; $[\alpha]_{\text{D}}^{25}$ -81° (CHCl₃, c = 0.53); ¹H NMR (400 MHz, CDCl₃) δ 1.11(s, 3 H, Me17), 1.12(s, 3 H, Me16), 1.60(s, 1 H, 1-OH), 1.69(s, 3 H, Me19), 1.87(ddd, J = 14.7, 11.0, 2.1 Hz, 1 H, H6b), 2.05(d, J = 5.1 Hz, 1 H, 13-OH), 2.08(d, J = 1.2, 3 H, Me18), 2.28(s, 3 H, 4-Ac), 2.29(m, 2 H, H14a, H14b), 2.47(d, J = 4.2 Hz, 1 H, 7-OH), 2.57(ddd, J = 14.7, 9.6, 6.7 Hz, 1 H, H6a), 3.86(d, J = 7.0 Hz, 1 H, H3), 4.16(d, J = 8.4 Hz, 1 H, H20b), 4.31(d, J = 8.4 Hz, 1 H, H20a), 4.44(ddd, J = 11.0, 6.7, 4.2 Hz, 1 H, H7), 4.70(br d, J = 5.9 Hz, 2 H, CHH'=CHCH₂O), 4.90(m, 1 H, H13), 4.97(dd, J = 9.6, 2.1 Hz, 1 H, H5), 5.32(dd, J = 10.4, 1.2 Hz, 1 H, CHH'=CHCH₂O), 5.42(dd, J = 17.2, 1.2 Hz, 1 H, CHH'=CHCH₂O), 5.63(d, J = 7.0 Hz, 1 H, H2), 5.98(ddt, J = 17.2, 10.4, 5.9 Hz, 1 H, CHH'=CHCH₂O), 6.16(s, 1 H, H10), 7.48(dd, J = 8.1, 7.5 Hz, 2 H, benzoate, m), 7.60(tt, J = 7.5, 1.2 Hz, 1 H, benzoate, p), 8.11(d, J = 8.1, 1.2 Hz, 2 H, benzoate, o) ppm; ¹³C NMR (75 MHz, CDCl₃) δ 9.1(Me(19)), 15.3(Me(18)), 20.7(4-Ac), 22.3, 26.7(Me16, Me17), 35.5(C(6)), 38.6(C(14)), 42.5(C(15)), 46.1(C(3)), 58.7(C(8)), 67.9(C(13)), 69.3(CH₂=CHCH₂O), 72.1, 75.0, 76.5(C(7), C(2), C(20)), 79.0, 79.1(C(1), C(10)), 80.9(C(4)), 84.5(C(5)), 119.6(CH₂=CHCH₂O), 128.8, 129.7,

130.3, 133.8 (benzoate), 131.4, 131.9 ($\text{CH}_2=\text{CHCH}_2\text{O}$, C(11)), 147.5 (C(12)), 155.4 (OC(O)O), 167.4 (benzoate), 170.9 (4-Ac), 204.7 (C(9)) ppm. Anal. Calcd for $\text{C}_{33}\text{H}_{40}\text{O}_{12}$: C, 63.05; H, 6.41. Found: C, 62.77; H, 6.48.

B. Selective acylation of a C(10) hydroxyl group using ZnCl_2 :

baccatin III. To a solution of 10-DAB (100 mg, 0.184 mmol) in THF (6 mL) at room temperature was added a mixture of acetic anhydride (6.5 mL) and ZnCl_2 /THF solution (0.5 M, 726 mL, 0.368 mmol, 2 equiv) under N_2 . The reaction mixture was stirred at room temperature for 4 h. Then the reaction mixture was diluted with EtOAc (100 mL), washed with saturated aqueous NaHCO_3 solution (40 mLx3), brine. The organic layer was dried over Na_2SO_4 , concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc:hexanes (1:1) as the eluent and dried *in vacuo* to give baccatin III as a colorless solid: yield, 100 mg (93%). mp 237-238 °C dec (ref 236-238 °C dec); $[\alpha]_{\text{Hg}} -63^\circ$ (CH_3OH , c = 0.45) (ref $[\alpha]_{\text{D}} -54^\circ$, CH_3OH); ^1H NMR (400 MHz, CDCl_3) δ 1.11(s, 6 H, Me16, Me17), 1.61(s, 1 H, 1-OH), 1.67(s, 3 H, Me19), 1.87(ddd, J = 14.7, 10.9, 2.1 Hz, 1 H, H6b), 2.05(d, J = 3.8 Hz, 1 H, 13-OH), 2.05(s, 3 H, Me18), 2.24(s, 3 H, 10-Ac), 2.28(s, 3 H, 4-Ac), 2.30(m, 2 H, H14a, H14b), 2.47(d, J = 4.2 Hz, 1 H, 7-OH), 2.57(ddd, J = 14.7, 9.4, 6.7 Hz, 1 H, H6a), 3.89(d, J = 7.0 Hz, 1 H, H3), 4.16(d, J = 8.4 Hz, 1 H, H20b), 4.31(d, J = 8.4 Hz, 1 H, H20a), 4.47(ddd, J = 10.9, 6.7, 4.2 Hz, 1 H, H7), 4.90(m, 1 H, H13), 4.99(dd, J = 9.4, 2.1 Hz, 1 H, H5), 5.63(d, J = 7.0 Hz, 1 H, H2), 6.33(s, 1 H, H10), 7.48(dd, J = 7.8, 7.8 Hz, 2 H, benzoate, m), 7.61(dd, J = 7.8, 7.4 Hz, 1 H, benzoate, p), 8.11(d, J = 7.4 Hz, 2 H, benzoate, o) ppm. ^{13}C NMR (100 MHz, CDCl_3) δ 9.4 (Me(19)),

15.6(Me(18)), 20.9(4-Ac, 10-Ac), 22.6, 27.0(Me16, Me17), 35.6(C(6)), 38.6(C(14)), 42.7(C(15)), 46.1(C(3)), 58.8(C(8)), 68.0(C(13)), 72.3, 75.0, 76.2, 76.4(C(7), C(2), C(10), C(20)), 79.1(C(1)), 80.9(C(4)), 84.5(C(5)), 128.6, 129.4, 130.1, 133.7(benzoate), 132.0(C(11)), 146.3(C(12)), 167.1(benzoate), 170.7, 171.3(10-Ac, 4-Ac), 204.1(C(9))ppm.

10-Chloroacetyl-10-DAB. To a solution of 10-DAB (116 mg, 0.21 mmol) in THF (3 mL) at room temperature was added a mixture of chloroacetic anhydride (2.8 g, 16.3 mmol, 78 equiv) and ZnCl_2 /THF solution (0.5 M, 0.85 mL, 0.42 mmol, 2 equiv) via a syringe under N_2 . The reaction mixture was stirred at room temperature for 5 h. The reaction mixture was poured into a mixture of EtOAc (200 mL) and saturated aqueous NaHCO_3 solution (100 mL). The organic layer was separated, and the aqueous layer was extracted with EtOAc (100 mLx3). The organic solution was combined, dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc:hexanes (1:1) as the eluent and dried overnight *in vacuo* to give 10-chloro-acetyl-10-DAB as a colorless solid: yield, 123 mg (93%). mp 231-233 °C dec; $[\alpha]_{\text{D}}^{25}$ -66° (EtOAc, $c = 0.45$); ^1H NMR (400 MHz, CDCl_3) δ 1.11(s, 3 H, Me17), 1.12(s, 3 H, Me16), 1.63(s, 1 H, 1-OH), 1.69(s, 3 H, Me19), 1.89(ddd, $J = 14.6, 10.9, 2.1$ Hz, 1 H, H6b), 2.07(d, $J = 5.2$ Hz, 1 H, 13-OH), 2.09(d, $J = 1.2, 3$ H, Me18), 2.12(d, $J = 4.5$ Hz, 1 H, 7-OH), 2.29(s, 3 H, 4-Ac), 2.30(m, 2 H, H14a, H14b), 2.58(ddd, $J = 14.6, 9.7, 6.7$ Hz, 1 H, H6a), 3.88(d, $J = 7.0$ Hz, 1 H, H3), 4.16(d, $J = 8.3$ Hz, 1 H, H20b), 4.27(br s, 2 H, ClCH_2), 4.31(d, $J = 8.3$ Hz, 1 H, H20a), 4.44(ddd, $J = 10.9, 6.7, 4.5$ Hz, 1 H, H7), 4.90(m, 1 H, H13), 4.98(dd, $J = 9.7, 2.1$ Hz, 1 H,

FSU 10293 012093

H5), 5.64(d, J = 7.0 Hz, 1 H, H2), 6.41(s, 1 H, H10), 7.49(dd, J = 7.9, 7.4 Hz, 2 H, benzoate, m), 7.61(tt, J = 7.4, 1.3 Hz, 1 H, benzoate, p), 8.11(d, J = 7.9, 1.3 Hz, 2 H, benzoate, o) ppm. ^{13}C NMR (75 MHz, CDCl_3) δ 9.3(Me(19)), 15.3(Me(18)), 20.6(4-Ac), 22.3, 26.7(Me16, Me17), 35.8(C(6)), 38.6(C(14)), 40.5(ClCH_2), 42.6(C(15)), 46.2(C(3)), 58.8(C(8)), 68.0(C(13)), 72.0, 75.0, 75.9(C(7), C(2), C(10), C(20)), 79.0(C(1)), 80.9(C(4)), 84.4(C(5)), 128.8, 129.7, 130.3, 133.9(benzoate), 131.8(C(11)), 147.1(C(12)), 167.4, 167.7($\text{ClCH}_2\text{C(O)O}$, benzoate), 171.0(4-Ac), 203.7(C(9))ppm. Anal. Calcd for $\text{C}_{31}\text{H}_{37}\text{ClO}_{11}$. H_2O : C, 58.26; H, 6.15. Found: C, 58.26; H, 6.07.

10-Propionyl-10-DAB. To a solution of 10-DAB (47 mg, 0.086 mmol) in THF (2 mL) at room temperature was added a mixture of propionic anhydride anhydride (4 mL) and ZnCl_2 / THF solution (0.5 M, 350 mL, 0.173 mmol, 2 equiv) under N_2 . The reaction mixture was stirred at room temperature for 14 h. Then the reaction mixture was diluted with EtOAc (150 mL), exhaustively washed with saturated aqueous NaHCO_3 solution (50 mLx3), brine. The organic layer was dried over Na_2SO_4 , concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:1) as the eluent and dried in vacuo to give 10-propionyl-10-DAB as a white solid: yield, 48 mg (93%). mp 212-213 °C dec; $[\alpha]_{\text{H}_g} -96^\circ$ (CHCl_3 , c = 0.78); ^1H NMR (400 MHz, CDCl_3) δ 1.11(s, 6 H, Me16, Me17), 1.24(t, J = 7.6 Hz, 3 H, CH_3CH_2), 1.60(s, 1 H, 1-OH), 1.67(s, 3 H, Me19), 1.87(ddd, J = 14.7, 10.9, 2.2 Hz, 1 H, H6b), 2.05(d, J = 5.1 Hz, 1 H, 13-OH), 2.06(d, J = 1.3 Hz, 3 H, Me18), 2.28(s, 3 H, 4-Ac), 2.30(d, J = 7.5 Hz, 2 H, H14a, H14b), 2.51(d, J = 4.1 Hz, 1 H, 7-OH), 2.55(q, J = 7.6 Hz, 2 H, CH_3CH_2),

2.57(ddd, J = 14.7, 9.5, 6.7 Hz, 1 H, H6a), 3.90(d, J = 6.9 Hz, 1 H, H3), 4.16(dd, J = 8.4, 0.8 Hz, 1 H, H20b), 4.31(d, J = 8.4 Hz, 1 H, H20a), 4.48(ddd, J = 10.9, 6.7, 4.1 Hz, 1 H, H7), 4.90(m, 1 H, H13), 4.99(dd, J = 9.5, 2.2 Hz, 1 H, H5), 5.63(d, J = 6.9 Hz, 1 H, H2), 6.34(s, 1 H, H10), 7.48(dd, J = 8.1, 7.4 Hz, 2 H, benzoate, m), 7.61(tt, J = 7.4, 1.3 Hz, 1 H, benzoate, p), 8.11(dd, J = 8.3, 1.3 Hz, 2 H, benzoate, o) ppm. ^{13}C NMR (75 MHz, CDCl_3) δ 8.8(CH_3CH_2), 9.2(Me(19)), 15.2(Me(18)), 20.7(4-Ac), 22.3, 26.8, 27.4(Me16, Me17, CH_3CH_2), 35.5(C(6)), 38.7(C(14)), 42.6(C(15)), 46.1(C(3)), 58.7(C(8)), 67.9(C(13)), 72.3, 75.1, 76.1, 76.5(C(7), C(2), C(10), C(20)), 79.1(C(1)), 80.9(C(4)), 84.5(C(5)), 128.7, 129.7, 130.3, 133.8(benzoate), 132.3(C(11)), 146.5(C(12)), 167.4(benzoate), 170.9, 174.9(4-Ac, 10-C(O)O), 204.6(C(9)) ppm. Anal. Calcd for $\text{C}_{32}\text{H}_{40}\text{O}_{11}$: C, 63.99; H, 6.71. Found: C, 63.81; H, 6.80.

C. Selective acylation of a C(10) hydroxyl group using CeCl_3 :

General procedure: To a solution of 10-DAB in THF (20 mL per mmol of 10-DAB) under N_2 was added CeCl_3 and the appropriate anhydride or pyrocarbonate (amounts specified in Table 1). The reaction mixture was stirred at 25 °C and monitored by TLC analysis. When that analysis indicated complete reaction (time specified in Table 1), the reaction mixture was diluted with EtOAc and washed three times with saturated aqueous sodium bicarbonate solution. The combined bicarbonate washings were extracted three times with EtOAc, the organic layers were combined and dried over sodium sulfate, and the solvent was evaporated. The crude product was purified by flash column chromatography. Further purification, if

necessary, was carried out by recrystallization from EtoAc/Hexane.

Table 1. CeCl_3 catalyzed acylation of 10-DAB

$10\text{-DAB} \xrightarrow[\text{THF, RT}]{(\text{RCO})_2\text{O, CeCl}_3} 10\text{-Acyl-10-DAB}$				
Entry	R (eq)	CeCl_3 (eq)	Time (hr)	Yield (%)
1	Me (10)	0.1	1.5	91
2	Pr (10)	0.1	3	100
3	iPr (10)	0.1	4.5	100
4	Ph (10)	0.1	21	94
5	cyclopropyl (10)	0.1	20.5	94
6	MeCH=CH (10)	0.1	20	91
7	$\text{CH}_2=\text{CHCH}_2\text{O}$ (5)	0.1	1	96
8	EtO (5)	0.1	3	99
9	MeO (5)	0.1	3	98
10	tBuO (10)	0.7	24	94
11	BnO (3)	0.7	1	98

10-butyryl-10-DAB. mp 145-149 °C; $[\alpha]_{\text{Hg}} -86.6^\circ$ (CHCl_3 , c = 1); ^1H NMR (500 MHz, CDCl_3) δ 8.13-8.11 (2H, m), 7.62 (1H, m), 7.51-7.48 (2H, m), 6.35 (1H, s), 5.64 (1H, d, J 7.0Hz), 4.99 (1H, d, J 7.7Hz), 4.90 (1H, m), 4.48 (1H, m), 4.31 (1H, d, J 8.3Hz), 4.18 (1H, d, J 8.3Hz), 3.91 (1H, d, J 7.0Hz), 2.60-2.42 (4H, m), 2.36-2.26 (2H, m), 2.28 (3H, s), 2.06 (3H, d, J 1.0Hz), 1.88 (1H, ddd, J 1.9, 10.9, 13.0Hz), 1.76 (2H, hex, J 7.4Hz), 1.68 (3H, s), 1.12 (6H, s) and 1.04 (3H, t, J 7.4Hz); ^{13}C NMR (100MHz, CDCl_3) δ 204.2, 173.9, 170.6, 167.1, 146.2, 133.7, 132.0, 130.1, 129.4, 128.6, 84.5, 88.9, 79.1, 76.5, 76.0, 75.0, 72.3, 68.0, 58.8, 46.2, 42.7, 38.7, 37.1, 36.2, 35.6, 30.6, 27.0, 22.6, 20.9,

18.4, 17.8, 15.5 and 9.4; Anal. Calcd. for $C_{33}H_{42}O_{11}$: C, 64.48; H, 6.89 Found: C, 63.67; H, 7.01.

10-isobutyryl-10-DAB. mp 143 °C; $[\alpha]_{\text{Hg}} -62.6^\circ$ (CHCl_3 , $c=0.075$); ^1H NMR (CDCl_3 , 500MHz): δ 8.12 (2H, d, J 7.3Hz), 7.62 (1H, m), 7.51-7.48 (2H, m), 6.33 (1H, s), 5.65 (1H, d, J 7.3Hz), 5.00 (1H, d, J 7.9Hz), 4.91 (1H, m), 4.48 (1H, ddd, J 4.3, 6.7, 11.0Hz), 4.31 (1H, d, J 8.6Hz), 4.18 (1H, d, J 8.6Hz), 3.91 (1H, d, J 7.3Hz), 2.74 (1H, pent, J 6.7Hz), 2.57 (1H, m), 2.51 (1H, d, J 4.3Hz), 2.31 (1H, m), 2.28 (3H, s), 2.06 (3H, s), 2.01 (1H, d, J 5.5Hz), 1.90 (1H, ddd, J 2.3, 11.0, 14.6Hz), 1.68 (3H, s), 1.60 (1H, s), 1.51 (3H, s), 1.33 (3H, d, J 6.7Hz), 1.26 (3H, d, J 6.7Hz), 1.13 (3H, s) and 1.12 (3H, s); ^{13}C NMR (100MHz, CDCl_3) δ 204.1, 177.2, 170.6, 167.1, 146.2, 133.7, 132.1, 130.1, 129.4, 128.6, 95.5, 84.5, 80.9, 79.1, 76.5, 75.8, 74.9, 72.3, 68.0, 58.8, 46.2, 42.7, 38.7, 35.6, 34.1, 27.0, 22.6, 20.9, 19.2, 18.7, 15.5 and 9.4; Anal. Calcd. for $C_{33}H_{42}O_{11} \cdot 0.5\text{H}_2\text{O}$: C, 64.48; H, 6.89 Found: C, 63.05; H, 6.70.

10-benzoyl-10-DAB. ^1H NMR (CDCl_3 , 500MHz): δ 8.15-8.11 (4H, m), 7.64-7.6 (2H, m), 7.52-7.48 (4H, m), 6.62 (1H, s), 5.7 (1H, d, J 7.1Hz), 5.02 (1H, d, J 7.7Hz), 4.94 (1H, m), 4.57 (1H, ddd, J 4.4, 7.1, 11.0Hz), 4.33 (1H, d, J 8.2Hz), 4.20 (1H, d, J 8.3Hz), 3.99 (1H, d, J 6.6Hz), 2.62 (1H, ddd, J 6.6, 9.3, 14.8), 2.55 (1H, d, J 4.4Hz), 2.35 (2H, m), 2.30 (3H, s), 2.13 (3H, d, J 1.1Hz), 2.03 (1H, d, J 4.9Hz), 1.91 (1H, ddd, J 2.2, 11.0, 13.2Hz), 1.71 (3H, s), 1.65 (1H, s), 1.25 (3H, s) and 1.21 (3H, s); ^{13}C NMR (100MHz, CDCl_3) δ 204.0, 170.7, 167.1, 166.5, 146.5, 133.7, 133.6, 132.0, 130.1, 129.9, 129.4, 129.3, 128.7, 128.5, 84.5, 80.9, 79.1, 76.5, 75.0, 72.4, 68.1, 58.8, 46.3, 42.8, 38.7, 35.8, 29.7, 27.2,

22.6, 21.2, 15.6 and 9.5; Anal. Calcd. for $C_{35}H_{40}O_{11}$: C, 66.66; H, 6.22 Found C, 66.46; H, 6.19.

10-trans crotonyl-10-DAB. 1H NMR ($CDCl_3$, 500MHz): δ 8.13 (2H, d, J 7.1Hz), 7.62 (1H, m), 7.51-7.48 (2H, m), 7.11 (1H, m), 6.42 (1H, s), 6.02 (1H, dq, J 1.7, 15.4Hz), 5.66 (1H, d, J 7.1Hz), 4.99 (1H, dd, J 2.0, 9.6Hz), 4.91 (1H, t, J 7.6Hz), 4.50 (1H, dd, J 7.1, 10.8Hz), 4.31 (1H, d, J 8.3Hz), 4.19 (1H, d, J 8.3Hz), 3.93 (1H, d, J 7.1Hz), 2.61-2.55 (2H, m), 2.33-2.31 (2H, m), 2.28 (3H, s), 2.07 (3H, d, J 1.5Hz), 1.95 (3H, dd, J 1.6, 6.8Hz), 1.89 (1H, ddd, J 2.3, 11.0, 13.4Hz), 1.68 (3H, s), 1.15 (3H, s) and 1.14 (3H, s); ^{13}C NMR (75MHz, $CDCl_3$) δ 212.4, 181.0, 170.8, 167.3, 166.5, 146.4, 133.8, 132.3, 130.2, 129.5, 128.7, 121.9, 116.0, 84.7, 84.6, 80.9, 79.2, 77.2, 75.9, 75.1, 72.4, 68.1, 58.8, 46.1, 42.7, 38.6, 35.6, 27.0, 20.9, 18.0, 15.4 and 9.3; Anal. Calcd. for $C_{33}H_{40}O_{11}$: C, 64.69; H, 6.58 Found C, 63.93; H, 6.61.

10-cyclopropanoyl-10-DAB. 1H ($CDCl_3$, 500MHz): δ 8.12 (2H, d, J 7.3Hz), 7.62 (1H, t, J 7.5Hz), 7.49 (2H, t, J 7.7Hz), 6.35 (1H, s), 5.65 (1H, d, J 7.0Hz), 4.99 (1H, app-d, J 8.2Hz), 4.91 (1H, m), 4.46 (1H, ddd, J 4.1, 6.8, 10.8Hz), 4.31 (1H, d, J 8.1Hz), 4.18 (1H, d, J 8.1Hz), 3.90 (1H, d, J 7.0Hz), 2.56 (1H, m), 2.51 (1H, d, J 4.1Hz), 2.31 (2H, m), 2.07 (3H, d, J 1.0Hz), 2.00 (1H, d, J 4.9Hz), 1.87 (1H, ddd, J 2.1, 10.8, 14.6Hz), 1.79 (1H, ddd, J 3.4, 7.9, 12.4Hz), 1.68 (3H, s), 1.60 (1H, s), 1.16-1.14 (2H, m), 1.13 (6H, s) and 1.01-0.97 (2H, m); ^{13}C NMR (100MHz, $CDCl_3$) δ 204.3, 175.2, 170.6, 167.1, 146.4, 133.7, 132.0, 130.1, 129.4, 128.6, 84.5, 80.9, 79.1, 76.5, 76.0, 74.9, 72.4, 68.0, 58.8, 46.2, 42.7, 38.6, 35.6, 34.0, 27.0, 25.6, 24.9, 22.6, 21.0, 15.6,

13.1, 9.4 and 9.1; Anal. Calcd. for $C_{33}H_{40}O_{11}$: C, 64.69; H, 6.58 Found: C, 64.47; H, 6.66.

10-Ethoxycarbonyl-10-DAB. mp 214-215 °C; $[\alpha]_{\text{Hg}} -81^\circ$ (CHCl_3 , $c = 0.35$); ^1H NMR (500 MHz, CDCl_3) δ 1.13(s, 3 H, Me17), 1.14(s, 3 H, Me16), 1.38(t, $J = 7.1$ Hz, 3 H, CH_3CH_2), 1.59(s, 1 H, 1-OH), 1.70(s, 3 H, Me19), 1.88(ddd, $J = 14.6, 10.5, 2.1$ Hz, 1 H, H6b), 2.00(d, $J = 5.0$ Hz, 1 H, 13-OH), 2.10(d, $J = 1.4$ Hz, 3 H, Me18), 2.28(s, 3 H, 4-Ac), 2.30(m, 2 H, H14a, H14b), 2.46(d, $J = 4.2$ Hz, 1 H, 7-OH), 2.57(ddd, $J = 14.6, 9.6, 6.7$ Hz, 1 H, H6a), 3.88(d, $J = 6.9$ Hz, 1 H, H3), 4.18(d, $J = 8.2$ Hz, 1 H, H20b), 4.31(d, $J = 8.2$ Hz, 1 H, H20a), 4.23-4.33(m, 2 H, CH_3CH_2), 4.44(ddd, $J = 10.5, 6.7, 4.2$ Hz, 1 H, H7), 4.90(m, 1 H, H13), 4.98(dd, $J = 9.6, 2.1$ Hz, 1 H, H5), 5.65(d, $J = 6.9$ Hz, 1 H, H2), 6.17(s, 1 H, H10), 7.48(dd, $J = 8.2, 7.3$ Hz, 2 H, benzoate, m), 7.60(tt, $J = 7.3, 1.4$ Hz, 1 H, benzoate, p), 8.11(d, $J = 8.2, 1.4$ Hz, 2 H, benzoate, o) ppm; ^{13}C NMR (75 MHz, CDCl_3) δ 9.2, 14.0, 15.5, 20.8, 22.4, 26.7, 35.4, 38.5, 42.4, 46.0, 58.6, 65.0, 67.7, 72.2, 74.9, 76.4, 78.7, 79.0, 80.6, 84.4, 128.7, 129.4, 130.1, 131.5, 133.7, 147.5, 155.4, 167.1, 170.8, 204.7 ppm.

10-Methoxycarbonyl-10-DAB. mp 218-219 °C; $[\alpha]_{\text{Hg}} -83^\circ$ (CHCl_3 , $c = 0.58$); ^1H NMR (500 MHz, CDCl_3) δ 1.12(s, 3 H, Me17), 1.13(s, 3 H, Me16), 1.59(s, 1 H, 1-OH), 1.70(s, 3 H, Me19), 1.88(ddd, $J = 14.7, 10.8, 1.8$ Hz, 1 H, H6b), 2.00(d, $J = 5.0$ Hz, 1 H, 13-OH), 2.10(d, $J = 1.4$ Hz, 3 H, Me18), 2.28(s, 3 H, 4-Ac), 2.30(m, 2 H, H14a, H14b), 2.40(d, $J = 4.1$ Hz, 1 H, 7-OH), 2.57(ddd, $J = 14.7, 9.7, 6.6$ Hz, 1 H, H6a), 3.87(d, $J = 6.9$ Hz, 1 H, H3), 3.88(s, 3 H, MeOC(O)), 4.18(d, $J = 8.4$ Hz, 1 H, H20b), 4.31(d, $J = 8.4$ Hz, 1 H, H20a), 4.44(ddd, $J = 10.8, 6.6, 4.1$ Hz, 1

[illegible]

D. Selective carbamoylation of a C(10) hydroxyl group:

General procedure for the Selective Carbamoylation of the C-10 Hydroxyl group of 10-DAB: A solution of 0.061 mmol (1.1 mol equiv) of the isocyanate in 2 mL of THF was added, under nitrogen, to a mixture of 10-DAB (30

mg, 0.055 mmol) and CuCl (5.5 mg, 0.055 mmol) at 0 °C. The mixture was stirred for the time indicated in Table 2. After this time the reaction was warmed to 25 °C and stirring was continued for the time indicated in Table 2. The reaction was quenched by the addition of saturated aqueous ammonium chloride solution, and the mixture was extracted three times with EtOAc. The combined organic layers were washed with saturated aqueous NaHCO₃ solution, dried over sodium sulfate, and the solvent was evaporated to yield a white solid. The product was purified by flash column chromatography using 2:1 EtOAc/hexane as the eluent.

Table 2. Carbamoylation of 10-DAB

10-DAB $\xrightarrow{\text{RNCO, CuCl, THF}}$ 10-Carbamoyl-10-DAB				
Entry	R (eq)	Temp (°C)	Time (hr)	Yield (%)
1	Et (1.1)	0	7.5	88
		rt	0.5	
2	allyl (1.1)	0	6	88
		rt	0.5	
3	Bu (1.1)	0	6.5	87
		rt	0.5	
4	Ph (1.1)	rt	3	94

10-ethylcarbamoyl-0-DAB. mp 241-243 °C; $[\alpha]_{\text{Hg}} -92.0^\circ$ (CHCl₃, c=0.5); ¹H NMR (400MHz, CDCl₃) δ 8.13 (2H, d, *J* 7.1Hz), 7.63 (1H, m), 7.52-7.48 (2H, m), 6.27 (1H, s), 5.63 (1H, d, *J* 6.9Hz), 5.01 (1H, dd, *J* 1.9, 9.6Hz), 4.97 (1H, m), 4.91 (1H, m), 4.50 (1H, ddd, *J* 3.7, 6.5, 10.5Hz), 4.31 (1H, d, *J* 8.3Hz), 4.17 (1H, d, *J* 8.3Hz), 3.88 (1H, d, *J* 7.0Hz), 3.32- 3.25 (2H, m), 3.10 (1H, d, *J* 3.7Hz), 2.56 (1H, ddd, *J* 6.8, 9.8, 14.8Hz), 2.31 (1H, m),

2.29 (3H, s), 2.09 (3H, s), 1.88 (1H, ddd, J 2.2, 11.0, 13.3Hz), 1.67 (3H, s), 1.60 (1H, s), 1.19 (3H, t, J 7.2Hz) and 1.10 (6H, s); Anal. Calcd. for $C_{32}H_{40}NO_{11}$: C, 62.43; H, 6.71 Found: C, 61.90; H, 6.77.

10-butylcarbamoyl-10-DAB. $[\alpha]_{\text{Hg}} -89.6^\circ$ (CHCl_3 , $c=0.25$); ^1H NMR (500MHz, CDCl_3) δ 8.12 (2H, d, J 7.3Hz), 7.61 (1H, m), 7.51- 7.45 (2H, m), 6.27 (1H, s), 5.64 (1H, d, J 6.7Hz), 5.00 (1H, d, J 8.0Hz), 4.91 (1H, m), 4.49 (1H, m), 4.31 (1H, d, J 8.5Hz), 4.19 (1H, d, J 8.5Hz), 3.89 (1H, d, J 6.7Hz), 3.25- 3.23 (2H, m), 3.04 (1H, m), 2.56 (1H, ddd, J 6.7, 9.7, 14.7Hz), 2.30 (1H, d, J 7.9Hz), 2.28 (3H, s), 2.09 (3H, s), 1.99 (1H, d, J 4.9Hz), 1.88 (1H, ddd, J 2.5, 11.0, 13.4Hz), 1.68 (3H, s), 1.59 (1H, s), 1.55 (2H, b), 1.42- 1.37 (2H, m), 1.11 (6H, s) and 0.95 (3H, t, J 7.6Hz); Anal. Calcd. for $C_{34}H_{44}NO_{11}$: C, 63.44; H, 7.05 Found: C, 62.64; H, 7.01.

10-phenylcarbamoyl-10-DAB. mp 178-180 $^\circ\text{C}$; $[\alpha]_{\text{Hg}} -93.0^\circ$ (CHCl_3 , $c=0.5$); ^1H NMR (400Hz, CDCl_3) δ 8.13 (2H, d, J 6.9Hz), 7.63 (1H, t, J 7.4Hz), 7.51 (2H, t, J 7.6Hz), 7.42 (1H, d, J 7.8Hz), 7.36- 7.32 (2H, m), 7.12 (1H, t, J 7.4Hz), 6.87 (1H, b), 6.38 (1H, s), 5.66 (1H, d, J 7.4Hz), 5.02 (1H, app d, J 7.8Hz), 5.93 (1H, m), 4.52 (1H, ddd, J 3.8, 6.5, 10.5Hz), 4.33 (1H, d, J 8.3Hz), 4.18 (1H, d, J 8.3Hz), 3.91 (1H, d, J 7.0Hz), 2.83 (1H, d, J 4.0Hz), 2.59 (1H, ddd, J 6.5, 9.4, 14.5Hz), 2.33 (1H, m), 2.29 (3H, s), 2.12 (3H, d, J 1.4Hz), 2.04 (1H, d, J 5.1Hz), 1.89 (1H, ddd, J 2.2, 11.0, 14.4Hz), 1.69 (3H, s), 1.62 (1H, s), 1.15 (3H, s) and 1.13 (3H, s).

10-allylcarbamoyl-10-DAB. mp 165-170 $^\circ\text{C}$; $[\alpha]_{\text{Hg}} -80.0^\circ$ (CHCl_3 , $c=0.25$); ^1H NMR (500MHz, CDCl_3) δ 8.12 (2H, d, J 7.3Hz), 7.62 (1H, m), 7.51- 7.48 (2H, m), 6.27 (1H,

s), 5.89 (1H, m), 5.62 (1H, d, J 6.7Hz), 5.31 (1H, s), 5.19 (1H, d, J 9.8Hz), 5.08 (1H, m), 5.00 (1H, d, J 7.9Hz), 4.90 (1H, m), 4.49 (1H, ddd, J, 3.7, 6.1, 10.4Hz), 4.31 (1H, d, J 8.5Hz), 4.17 (1H, d, J 8.5Hz), 3.88- 3.86 (2H, m), 3.03 (1H, d, J 3.7Hz), 2.55 (1H, ddd, J 6.7, 9.8, 15.9Hz), 2.30 (1H, m), 2.29 (3H, s), 2.08 (3H, s), 2.06 (1H, app d, J 4.9Hz), 1.87 (1H, ddd, J 1.8, 11.0, 14.0Hz), 1.67 (3H, s), 1.58 (1H, s) and 1.09 (6H, s); Anal. Calcd. for $C_{33}H_{40}NO_{11}$: C, 63.15; H, 6.58 Found: C, 61.73; H, 6.45.

E. Selective silylation of a C(10) hydroxyl group:

10-TMS-10-DAB. To a solution of 10-DAB (100 mg, 0.18 mmol) in THF (10 mL) at 0 °C was slowly added N, O-bis(trimethylsilyl)trifluoroacetamide (1.0 mL, 3.7 mmol, 20 equiv) under N_2 . The reaction mixture was stirred at 0 °C for 5 h. EtOAc (20 mL) was added, and the solution was filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:1) as the eluent and dried in vacuo overnight to give 10-TMS-10-DAB as a white solid: yield, 103 mg (91%). mp 189-191 °C; $[\alpha]_{Hg} -70^\circ$ ($CHCl_3$, c = 0.55); 1H NMR (400 MHz, $CDCl_3$) δ 0.18(s, 9 H, Me_3Si), 1.06(s, 3 H, Me17), 1.16(s, 3 H, Me16), 1.31(d, J = 8.6 Hz, 1 H, 7-OH), 1.56(s, 1 H, 1-OH), 1.68(s, 3 H, Me19), 1.79(ddd, J = 14.4, 11.1, 2.1 Hz, 1 H, H6b), 1.97(d, J = 4.9 Hz, 1 H, 13-OH), 2.03(d, J = 1.3 Hz, 3 H, Me18), 2.27(m, 2 H, H14a, H14b), 2.28(s, 3 H, 4-Ac), 2.58(ddd, J = 14.4, 9.6, 7.5 Hz, 1 H, H6a), 4.01(d, J = 7.2 Hz, 1 H, H3), 4.16(d, J = 8.2 Hz, 1 H, H20b), 4.25(ddd, J = 11.1, 8.6, 7.5 Hz, 1 H, H7), 4.30(d, J = 8.2 Hz, 1 H, H20a), 4.84(m, 1 H, H13), 4.97(dd, J = 9.6, 2.1 Hz, 1 H, H5),

5.27(s, 1 H, H10), 5.64(d, J = 7.2 Hz, 1 H, H2), 7.47(dd, J = 8.2, 7.5 Hz, 2 H, benzoate, m), 7.60(tt, J = 7.5, 1.2 Hz, 1 H, benzoate, p), 8.11(dd, J = 8.2, 1.2 Hz, 2 H, benzoate, o) ppm. ^{13}C NMR (75 MHz, CDCl_3) δ 0.2(Me_3Si), 9.7(Me(19)), 14.4(Me(18)), 19.6(4-Ac), 22.4, 26.6(Me16, Me17), 37.1(C(6)), 38.6(C(14)), 42.6(C(15)), 47.2(C(3)), 57.8(C(8)), 68.0(C(13)), 72.0, 75.1, 76.1, 76.8(C(7), C(2), C(10), C(20)), 78.9(C(1)), 81.2(C(4)), 84.3(C(5)), 128.8, 130.3, 133.7(benzoate), 137.0(C(11)), 139.0(C(12)), 167.4(benzoate), 171.0(4-Ac), 209.5(C(9)) ppm. Anal. Calcd for $\text{C}_{32}\text{H}_{44}\text{O}_{10}\text{Si}$. $1/2\text{H}_2\text{O}$: C, 61.42; H, 7.25. Found: C, 61.61; H, 7.12.

10-TES-10-DAB. To a solution of 10-DAB (85 mg, 0.16 mmol) in THF (3 mL) at 0 °C was slowly added N, O-bis(triethylsilyl)trifluoroacetamide (484 mL, 1.56 mmol, 10 equiv), and a catalytic amount of LiHMDS/THF solution (1 M, 5 mL, 0.005 mmol), respectively, under N_2 . The reaction mixture was stirred at 0 °C for 5 min. EtOAc (10 mL) was added, and the solution was filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:2) as the eluent and dried in vacuo overnight to give the 10-TES-10-DAB as a white solid: yield, 98 mg (95 %). mp 234-235 °C dec; $[\alpha]_{\text{D}}^{25}$ -69° (CHCl_3 , c = 0.95); IR 3690, 2958, 1714, 1602 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ 0.68(m, 6 H, $(\text{CH}_3\text{CH}_2)_3\text{Si}$), 1.00(t, J = 7.9, 9 H, $(\text{CH}_3\text{CH}_2)_3\text{Si}$), 1.08(s, 3 H, Me17), 1.19(s, 3 H, Me16), 1.29(d, J = 8.4 Hz, 1 H, 7-OH), 1.55(s, 1 H, 1-OH), 1.69(s, 3 H, Me19), 1.79(ddd, J = 14.4, 11.0, 2.0 Hz, 1 H, H6b), 1.92(d, J = 5.0 Hz, 1 H, 13-OH), 2.03(d, J = 1.0 Hz, 3 H, Me18), 2.27(s, 3 H, 4-Ac), 2.29(m, 2 H, H14a, H14b), 2.59(ddd, J = 14.4, 9.5,

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6.7 Hz, 1 H, H6a), 4.02(d, J = 7.2 Hz, 1 H, H3), 4.18(d, J = 8.5 Hz, 1 H, H20b), 4.23(ddd, J = 11.0, 8.4, 6.7 Hz, 1 H, H7), 4.30(d, J = 8.5 Hz, 1 H, H20a), 4.86(m, 1 H, H13), 4.97(dd, J = 9.5, 2.0 Hz, 1 H, H5), 5.28(s, 1 H, H10), 5.66(d, J = 7.2 Hz, 1 H, H2), 7.47(dd, J = 7.9, 7.9 Hz, 2 H, benzoate, m), 7.59(tt, J = 7.9, 1.0 Hz, 1 H, benzoate, p), 8.11(dd, J = 7.9, 1.0 Hz, 2 H, benzoate, o)ppm. ¹³C NMR (75 MHz, CDCl₃) δ 4.9, 6.5(TES), 9.7(Me(19)), 14.3(Me(18)), 19.6(4-Ac), 22.4, 26.6(Me16, Me17), 37.1(C(6)), 38.6(C(14)), 42.6(C(15)), 47.3(C(3)), 57.9(C(8)), 67.9(C(13)), 71.9, 75.1, 76.1, 76.7(C(7), C(2), C(10), C(20)), 78.9(C(1)), 81.2(C(4)), 84.3(C(5)), 128.7, 129.9, 130.3, 133.7(benzoate), 137.0(C(11)), 138.8(C(12)), 167.4(benzoate), 171.0(4-Ac), 209.5(C(9)) ppm. Anal. Calcd for C₃₅H₅₀O₁₀Si. H₂O: C, 62.11; H, 7.74. Found: C, 62.45; H, 7.74.

Example 2

General Procedure for the Preparation of 7-silyl-10-TES-10-DAB. To a solution of 7-triethylsilyl-10-DAB, 7-t-butyldimethylsilyl-10-DAB, or 7-dimethylisopropylsilyl-10-DAB in THF at 0 °C was slowly added N, O-bis(triethylsilyl)trifluoroacetamide (5 equiv), and a catalytic amount of LiHMDS/THF solution (5 mol %), respectively, under N₂. The reaction mixture was stirred at 0 °C for 15 min. EtOAc (10 mL) was added, and the solution was filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:2) as the eluent and dried in vacuo overnight to give, respectively, 7,10-bis(triethylsilyl)-10-DAB (95% yield), 7-t-butyldimethylsilyl-10-

triethylsilyl-10-DAB (98% yield), or 7-dimethylisopropylsilyl-10-triethylsilyl-10-DAB (94% yield).

7-Dimethylphenylsilyl-10-TBS-10-DAB. To a solution of 7-dimethylphenylsilyl-10-DAB (35 mg, 0.052 mmol) in THF (2 mL) at 0 °C was added N,O-bis(t-butylldimethylsilyl)trifluoroacetamide (337 µL, 1.09 mmol, 20 equiv), and a catalytic amount of LiHMDS/THF solution (1 M, 6 µL, 0.006 mmol), respectively, under N₂. The reaction mixture was stirred at 0 °C for 4 h, then warmed to room temperature for an additional 4 h. EtOAc (10 mL) was added, and the solution was filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:2) as the eluent and dried in vacuo overnight to give 39 mg (92% yield) of 7-dimethylphenylsilyl-10-t-butylldimethylsilyl-10-DAB.

Example 3

Selective silylation of a C(7) hydroxyl group

7-TBS-10-DAB. To a mixture of 10-DAB (38 mg, 0.070 mmol), imidazole (190 mg, 2.79 mmol, 40 equiv), and tert-butylldimethylsilyl chloride (210 mg, 1.40 mmol, 20 equiv) was added DMF (0.1 mL) at room temperature under N₂. The reaction mixture was vigorously stirred at room temperature for 24 h. EtOAc (20 mL) was added, and the solution was filtered through a short column of silica gel. The silica gel was washed with EtOAc (200 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using 10% EtOAc-CH₂Cl₂ as the eluent and dried in vacuo overnight to give 7-TBS-10-DAB as a white solid: yield,

41 mg (90%). mp 222-223 °C; $[\alpha]_{\text{Hg}} -51^\circ$ (CHCl_3 , $c = 0.36$); ^1H NMR (400 MHz, CDCl_3) δ 0.05, 0.06 (2 s, 6 H, Me_2Si), 0.83 (s, 9 H, Me_3C), 1.09 (s, 6 H, Me16, Me17), 1.57 (s, 1 H, 1-OH), 1.75 (s, 3 H, Me19), 1.87 (ddd, $J = 14.4, 10.6, 2.0$ Hz, 1 H, H6b), 2.01 (d, $J = 5.0$ Hz, 1 H, 13-OH), 2.09 (d, $J = 1.3$, 3 H, Me18), 2.28 (m, 2 H, H14a, H14b), 2.29 (s, 3 H, 4-Ac), 2.46 (ddd, $J = 14.4, 9.6, 6.7$ Hz, 1 H, H6a), 3.96 (d, $J = 6.9$ Hz, 1 H, H3), 4.16 (d, $J = 8.3$ Hz, 1 H, H20b), 4.24 (d, $J = 2.2$ Hz, 1 H, 10-OH), 4.31 (d, $J = 8.3$ Hz, 1 H, H20a), 4.38 (dd, $J = 10.6, 6.7$ Hz, 1 H, H7), 4.88 (m, 1 H, H13), 4.96 (dd, $J = 9.6, 2.0$ Hz, 1 H, H5), 5.15 (d, $J = 2.0$ Hz, 1 H, H10), 5.60 (d, $J = 6.9$ Hz, 1 H, H2), 7.47 (dd, $J = 8.1, 7.5$ Hz, 2 H, benzoate, m), 7.60 (tt, $J = 7.5, 1.3$ Hz, 1 H, benzoate, p), 8.10 (d, $J = 8.1, 1.3$ Hz, 2 H, benzoate, o) ppm. ^{13}C NMR (75 MHz, CDCl_3) δ -5.8, -3.8 (Me_2Si), 9.7 (Me(19)), 14.8 (Me(18)), 17.6 (Me_3C), 19.3 (4-Ac), 22.4, 26.7 (Me16, Me17), 25.4 (Me_3C), 37.4 (C(6)), 38.7 (C(14)), 42.7 (C(15)), 47.0 (C(3)), 58.0 (C(8)), 68.0 (C(13)), 73.1, 74.7, 75.0 (C(7), C(2), C(10), C(20)), 78.9 (C(1)), 80.9 (C(4)), 84.3 (C(5)), 128.8, 129.8, 130.3, 133.8 (benzoate), 135.7 (C(11)), 141.9 (C(12)), 167.4 (benzoate), 171.2 (4-Ac), 210.8 (C(9)) ppm. Anal. Calcd for $\text{C}_{35}\text{H}_{50}\text{O}_{10}\text{Si}$: C, 63.80; H, 7.65. Found: C, 63.72; H, 7.70.

7-Dimethylphenylsilyl-10-DAB. To a THF (3 mL) solution of 10-DAB (54 mg, 0.099 mmol) at -20 °C was added pyridine (0.6 mL), dimethylphenylsilyl chloride (250 mL, 1.49 mmol, 15 equiv) under N_2 . The reaction mixture was stirred at -20 °C for 2 h. EtOAc (10 mL) and saturated NaHCO_3 aqueous solution (0.5 mL) was added, and the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced

pressure. The residue was purified by flash column chromatography using EtOAc: CH₂Cl₂ (1:10) as the eluent and dried overnight in vacuo to give 7-dimethylphenylsilyl-10-DAB as a white solid: yield, 62 mg (92%). mp 219-220 °C; $[\alpha]_{\text{Hg}} -28^\circ$ (CHCl₃, c = 0.27); ¹H NMR (400 MHz, CDCl₃) δ 0.35, 0.37(2 s, 6 H, Me₂Si), 1.05(s, 3 H, Me17), 1.06(s, 3 H, Me16), 1.54(s, 1 H, 1-OH), 1.73(d, J = 1.1, 3 H, Me18), 1.76(s, 3 H, Me19), 1.90(ddd, J = 14.4, 10.6, 2.1 Hz, 1 H, H6b), 1.93(d, J = 5.0 Hz, 1 H, 13-OH), 2.23(m, 2 H, H14a, H14b), 2.25(s, 3 H, 4-Ac), 2.43(ddd, J = 14.4, 9.6, 6.8 Hz, 1 H, H6a), 3.86(d, J = 7.0 Hz, 1 H, H3), 4.10(d, J = 2.1 Hz, 1 H, 10-OH), 4.16(d, J = 8.3 Hz, 1 H, H20b), 4.28(d, J = 8.3 Hz, 1 H, H20a), 4.31(dd, J = 10.6, 6.8 Hz, 1 H, H7), 4.81(m, 1 H, H13), 4.84(d, J = 2.1 Hz, 1 H, H10), 4.90(dd, J = 9.6, 2.1 Hz, 1 H, H5), 5.59(d, J = 7.0 Hz, 1 H, H2), 7.41, 7.53(2 m, 5 H, C₆H₅), 7.46(dd, J = 8.0, 7.5 Hz, 2 H, benzoate, m), 7.55(tt, J = 7.5, 1.2 Hz, 1 H, benzoate, p), 8.09(d, J = 8.0, 1.2 Hz, 2 H, benzoate, o) ppm. ¹³C NMR (75 MHz, CDCl₃) δ -1.8, -1.1(Me₂Si), 9.8(Me(19)), 14.4(Me(18)), 19.4(4-Ac), 22.3, 26.7(Me16, Me17), 37.2(C(6)), 38.6(C(14)), 42.6(C(15)), 46.7(C(3)), 58.0(C(8)), 68.0(C(13)), 73.2, 74.7, 75.0(C(7), C(2), C(10), C(20)), 78.8(C(1)), 80.8(C(4)), 84.3(C(5)), 128.3, 128.8, 129.8, 130.2, 130.3, 133.65, 133.74(PhSi, benzoate), 135.4(C(11)), 142.1(C(12)), 167.4(benzoate), 171.0(4-Ac), 210.9(C(9))ppm. Anal. Calcd for C₃₇H₄₆O₁₀Si. 1/2H₂O: C, 64.61; H, 6.89. Found: C, 64.72; H, 6.81.

7-Dimethylisopropylsilyl-10-DAB. To a solution of 10-DAB (97 mg, 0.18 mmol) in pyridine (1 mL) at -10 °C was added dimethylisopropylsilyl chloride (580 mL, 3.57 mmol, 20 equiv) under N₂. The reaction mixture was stirred at -10 °C for 3 h. EtOAc (10 mL) was added, and the solution

was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (150 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:2) as the eluent to give 7-dimethylisopropyl-10-DAB as a white solid: yield, 107 mg (93%). mp 229-230 °C; $[\alpha]_{\text{D}}^{25}$ -56° (CHCl₃, c = 0.62); ¹H NMR (400 MHz, CDCl₃) δ 0.05, 0.06 (2 s, 6 H, Me₂Si), 0.70 (m, 1 H, CHSi), 0.90, 0.92 (2 dd, J = 7.4, 1.7, 6 H, Me₂CH), 1.09 (s, 6 H, Me16, Me17), 1.56 (s, 1 H, 1-OH), 1.74 (s, 3 H, Me19), 1.89 (ddd, J = 14.4, 10.6, 2.1 Hz, 1 H, H6b), 1.99 (d, J = 5.0 Hz, 1 H, 13-OH), 2.09 (d, J = 1.4, 3 H, Me18), 2.28 (d, J = 7.9, 2 H, H14a, H14b), 2.29 (s, 3 H, 4-Ac), 2.44 (ddd, J = 14.4, 9.7, 6.7 Hz, 1 H, H6a), 3.96 (d, J = 7.3 Hz, 1 H, H3), 4.17 (d, J = 8.3 Hz, 1 H, H20b), 4.24 (d, J = 2.2 Hz, 1 H, 10-OH), 4.31 (d, J = 8.3 Hz, 1 H, H20a), 4.38 (dd, J = 10.6, 6.7 Hz, 1 H, H7), 4.85 (m, 1 H, H13), 4.95 (dd, J = 9.7, 2.1 Hz, 1 H, H5), 5.15 (d, J = 2.2 Hz, 1 H, H10), 5.61 (d, J = 7.3 Hz, 1 H, H2), 7.47 (dd, J = 8.2, 7.5 Hz, 2 H, benzoate, m), 7.60 (tt, J = 7.5, 1.4 Hz, 1 H, benzoate, p), 8.10 (d, J = 8.2, 1.4 Hz, 2 H, benzoate, o) ppm. ¹³C NMR (75 MHz, CDCl₃) δ -4.6, -3.3 (Me₂Si), 9.7 (Me(19)), 14.8, 14.9 (CHSi, Me(18)), 16.4, 16.5 (Me₂CH), 19.4 (4-Ac), 22.4, 26.7 (Me16, Me17), 37.3 (C(6)), 38.7 (C(14)), 42.7 (C(15)), 47.0 (C(3)), 58.0 (C(8)), 68.0 (C(13)), 73.1, 74.7, 75.0 (C(7), C(2), C(10), C(20)), 78.9 (C(1)), 80.9 (C(4)), 84.3 (C(5)), 128.8, 129.8, 130.3, 133.7 (benzoate), 135.7 (C(11)), 142.0 (C(12)), 167.4 (benzoate), 171.1 (4-Ac), 210.8 (C(9)) ppm. Anal. Calcd for C₃₄H₄₈O₁₀Si. H₂O: C, 61.61; H, 7.60. Found: C, 61.30; H, 7.35.

7-Tribenzylsilyl-10-DAB. To a mixture of 10-DAB (62 mg, 0.11 mmol), imidazole (280 mg, 4.11 mmol, 36 equiv)

and tribenzylsilyl chloride (364 mg, 1.14 mmol, 10 equiv) was added DMF (0.4 mL) under N₂. The reaction mixture was stirred at room temperature for 3 h. EtOAc (30 mL) was added, and the solution was filtered through a short column of silica gel. The silica gel was washed with EtOAc (150 mL), and the solution was concentrated under reduced pressure. The residue was purified twice by flash column chromatography, first time using EtOAc: hexanes (1:2) as the eluent, second time using EtOAc: CH₂Cl₂ as the eluent, and dried overnight in vacuo to give the 7-tribenzylsilyl-10-DAB as a white solid: yield, 88 mg (91%). mp 161-163 °C; IR 3690, 2928, 2890, 1712, 1600 cm⁻¹; [α]_D²⁵ -46° (CHCl₃, c = 0.46); ¹H NMR (400 MHz, CDCl₃) δ 1.10(s, 3 H, Me17, Me16), 1.56(s, 1 H, 1-OH), 1.71(ddd, J = 14.2, 10.9, 2.0 Hz, 1 H, H6b), 1.74(s, 3 H, Me19), 2.00(d, J = 5.1 Hz, 1 H, 13-OH), 2.07(ddd, J = 14.2, 9.6, 6.6 Hz, 1 H, H6a), 2.10(d, J = 1.2, 3 H, Me18), 2.12(s, 6 H, (PhCH₂)₃Si), 2.27(d, J = 7.5 Hz, 2 H, H14a, H14b), 2.27(s, 3 H, 4-Ac), 3.99(d, J = 7.0 Hz, 1 H, H3), 4.16(d, J = 8.5 Hz, 1 H, H20b), 4.18(d, J = 2.2 Hz, 1 H, 10-OH), 4.28(d, J = 8.5 Hz, 1 H, H20a), 4.58(dd, J = 10.9, 6.6 Hz, 1 H, H7), 4.81(dd, J = 9.6, 2.0 Hz, 1 H, H5), 4.89(m, 1 H, H13), 5.21(d, J = 2.2 Hz, 1 H, H10), 5.61(d, J = 7.0 Hz, 1 H, H2), 6.93, 7.09, 7.20(3 m, 15 H, (PhCH₂)₃Si), 7.48(dd, J = 8.1, 7.5 Hz, 2 H, benzoate, m), 7.61(tt, J = 7.5, 1.3 Hz, 1 H, benzoate, p), 8.10(d, J = 8.1, 1.3 Hz, 2 H, benzoate, o) ppm. ¹³C NMR (75 MHz, CDCl₃) δ 9.9(Me(19)), 15.0(Me(18)), 19.5(4-Ac), 22.4, 26.7(Me16, Me17), 23.6(Si(CH₂Ph)₃), 36.9(C(6)), 38.7(C(14)), 42.7(C(15)), 46.8(C(3)), 58.0(C(8)), 68.0(C(13)), 74.4, 74.9, 75.0(C(7), C(2), C(10), C(20)), 78.8(C(1)), 80.8(C(4)), 84.1(C(5)), 124.9, 128.7, 128.8, 129.1, 129.8, 130.3, 133.8, 137.8(Si(CH₂Ph)₃, benzoate), 135.5(C(11)), 142.2(C(12)), 167.4(benzoate), 170.9(4-Ac),

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210.8(C(9))ppm. Anal. Calcd for $C_{50}H_{56}O_{10}Si \cdot 1/2H_2O$: C, 70.32; H, 6.73. Found: C, 70.11; H, 6.57.

Example 4

Selective acylation of 10-acyl-10-DAB

10-Alloc-7-*p*-Nitrobenzyloxycarbonyl-10-DAB. To a mixture of 10-alloc-10-DAB (33 mg, 0.053 mmol) and DMAP (19.3 mg, 0.16 mmol, 3 equiv) in dichloromethane (4 mL) at 0 °C was added a dichloromethane solution (1 mL) of *p*-nitrobenzyl chloroformate (23 mL, 0.11 mmol, 2 equiv) under N_2 . The reaction mixture was stirred at 0 °C for 4 h. EtOAc (10 mL) was added, the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:2) as the eluent and dried overnight in vacuo to give 10-alloc-7-*p*-nitrobenzyloxycarbonyl-10-DAB as a colorless solid: yield, 34 mg (92%).

7-Benzyloxycarbonyl baccatin III. To a stirred solution of baccatin III (100 mg, 0.168 mmol) in methylene chloride under N_2 at room temperature was added 4-dimethylaminopyridine (204 mg, 1.68 mmol) followed by addition of benzyl chloroformate (240 mL, 1.68 mmol). The reaction mixture was stirred at room temperature and the progress of the reaction was monitored by TLC. After about 4 h the reaction was complete. The mixture was diluted with EtOAc (10 mL) and was transferred to a separatory funnel containing 50 mL of a 50% EtOAc/Hexanes. The mixture was washed with saturated sodium bicarbonate and the organic layer was separated. The aqueous layer was washed with 20 mL of 50% EtOAc/Hexanes. The combined organic layers were washed with brine, dried

over MgSO_4 , and concentrated under reduced pressure. The crude product was passed through a short column to give 115 mg (95%) of a white solid m.p. $245-248^\circ\text{C}$; $[\alpha]^{25}_{\text{D}}$ -60.5° c ($C=0.007$, CHCl_3). ^1H NMR (CDCl_3 , 400 MHz) δ 8.10(d, $J=9.6$ Hz, 2H, o-benzoate), 7.60-6.8 (m, 8H, benzoate, Bn), 6.45(s, 1H, H10), 5.63(d, $J=6.9$ Hz, 1H, H2b), 5.56 (dd, $J=10.6, 7.2$ Hz, 1H H7), 5.56(dd, $J=18.5, 12.0$ Hz, 2H, Bn), 4.97(d, $J=10.6$, 1H, H5), 4.87(m, 1H, H13), 4.31(d, $J=10.5$, 1H, H20a), 4.15 (d, $J=10.5$, 1H, H20b), 4.02(d, $J=6.9$, 1H, H3), 2.61(m, 1H, H6a), 2.30(m, 2H, H14's), 2.29(s, 3H, 4Ac), 2.18(s, 3H, 10Ac), 2.15(br s, 3H, Me18), 2.08(d, $J=5.2$ Hz, 13OH), 1.94(m, 1H, 6b), 1.79(s, 3H, Me19), 1.58(s, 1H, 1OH), 1.14(s, 3H, Me16), 1.09(s, 3H, Me17).

7-Allyloxycarbonyl baccatin III. To a stirred solution of baccatin III (30 mg, 0.051 mmol) in methylene chloride (1 mL) under N_2 at room temperature, was added 4-dimethylaminopyridine (62.3 mg, 0.51 mmol) followed by addition of allyl chloroformate (54 mL, 0.51 mmol). The reaction mixture was stirred at room temperature and the progress of the reaction was followed by TLC. After about 1.5 h the reaction was complete. The mixture was diluted by EtOAc (5 mL) and was transferred to a separatory funnel containing 50 mL of a 50% EtOAc/Hexanes. The mixture was washed with saturated sodium bicarbonate and the organic layer was separated. The aqueous layer was washed with 10 mL of 50% EtOAc/Hexanes, the combined organic layers were washed with brine, dried over MgSO_4 , and concentrated under reduced pressure. The crude product was passed through a short column to give 33.1 mg (97%) of a white solid m.p. $239-244^\circ\text{C}$; $[\alpha]^{25}_{\text{D}}$ -61.5° c (0.01, CHCl_3) ^1H NMR (CDCl_3 , 500 MHz) δ 8.12(d, $J=8.3$ Hz, 2H, o--benzoate), 7.66-7.45 (m,

3H, benzoate), 6.43(s, 1H, H10), 5.97(m, 1H, int. allyl), 5.64(d, J=7.0 Hz, 1H, H2b), 5.54 (dd, J=10.5, 7.0 Hz, 1H, H7), 5.28(m, 2H, ext. allyl), 4.97(d, J=9.6 Hz, 1H, H5), 4.87(m, 1H, H13), 4.67(m, 2H, CH2allyl), 4.31(d, J=8.5 Hz, 1H, H20a), 4.17(d, J=8.5, 1H, H20b), 4.02(d, J=7.0, 1H, H3), 2.64(m, 1H, H6a), 2.30(d, J=8.0 Hz, 2H, H14's), 2.29(s, 3H, 4Ac), 2.16(s, 3H, 10Ac), 2.15(br s, 3H, Me18), 2.01(d, J=5 Hz, 13OH), 1.96(m, 1H, 6b), 1.81(s, 3H, Me19), 1.58(s, 1H, 10H), 1.15(s, 3H, Me16), 1.02(s, 3H, Me17).

Example 5

Selective ketalization of 10-acyl-10-DAB

7-MOP baccatin III. To a solution of baccatin III (101 mg, 0.172 mmol) in THF (8 mL) at -20 °C was added 2-methoxypropene (0.66 mL, 6.89 mmol, 40 equiv), followed by the addition of a catalytic amount of toluenesulfonic acid (0.1 M solution in THF, 43 μ L, 0.004 mmol, 0.025 equiv) under N₂. The reaction mixture was stirred at -20 °C for 3 h. TLC analysis indicated complete consumption of the starting material and the formation of desired product as the only major product. Triethylamine (0.5 mL) was added, and the solution was warmed to room temperature, diluted with EtOAc (100 mL), washed with saturated aqueous NaHCO₃ solution, dried over Na₂SO₄, and concentrated under reduced pressure. The residue was dried in vacuo overnight to give 112 mg (99%) of crude product. Recrystallization of the crude product from EtOAc/hexanes gave 105 mg (93%) of 7-MOP baccatin III as a white crystal, mp 181-183 °C; ¹H NMR (500 MHz, C₆D₆) δ 1.01 (s, 3 H, Me17), 1.11(br s, 1 H, 13-OH), 1.28(s, 3 H, Me16), 1.39, 1.78(2 s, 6 H, Me₂CO), 1.62(s, 1 H, 1-OH), 1.78(s, 3 H, 10-Ac), 1.92(s, 3 H, 4-Ac), 2.09(s, 3 H, Me18), 2.12(s, 3 H, Me19), 2.14(ddd, J = 15.0, 10.9, 2.2

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Hz, 1 H, H6b), 2.18(dd, J = 15.6, 9.4 Hz, 1 H, H14b), 2.31(dd, J = 15.6, 7.0 Hz, 1 H, H14b), 2.97(s, 3 H, MeO), 3.15(ddd, J = 15.0, 9.9, 6.7 Hz, 1 H, H6a), 4.08(d, J = 7.0 Hz, 1 H, H3), 4.24(m, 1 H, H13), 4.33(d, J = 8.3 Hz, 1 H, H20b), 4.41(d, J = 8.3 Hz, 1 H, H20a), 4.78(dd, J = 10.9, 6.7 Hz, 1 H, H7), 4.97(dd, J = 9.9, 2.2 Hz, 1 H, H5), 5.95(d, J = 7.0 Hz, 1 H, H2), 6.79(s, 1 H, H10), 7.15(m, 3 H, benzoate, m, p), 8.28(d, J = 8.0 Hz, 2 H, benzoate, o) ppm; Anal. Calcd for $C_{35}H_{46}O_{12}$: C, 63.82; H, 7.04. Found: C, 63.72; H, 7.07.

Example 6

Selective acylation of 10-silyl-10-DAB

7-Acetyl-10-TES-10-DAB. To a stirred solution of 10-TES-10-DAB (65 mg, 0.098 mmol) in dichloromethane (4 ml) at 0 °C under N_2 , was added DMAP (36 mg, 0.296 mmol, 3 equiv), followed by addition of acetic anhydride (14 mL, 0.148 mmol, 1.5 equiv). The reaction mixture was stirred at 0 °C for 4.5 hrs and the TLC analysis indicated the complete consumption of starting material. The reaction mixture was then filtered through a short pad of silica gel, the silica gel was washed with EtOAc (100 mL) and the solution was concentrated under reduced pressure. The residue was purified by flash chromatography using EtOAc:hexanes (1:2) as the eluent and dried in vacuo overnight to give the 7-acetyl-10-TES-10-DAB: yield, 65.7 mg (95%). 1H NMR ($CDCl_3$, 400 MHz), δ 0.60(m, 6 H, $(CH_3CH_2)_3Si$), 0.97(t, J = 7.9 Hz, 9 H, $(CH_3CH_2)_3Si$), 1.05(s, 3 H, Me17), 1.18(s, 3 H, Me16), 1.56(s, 1 H, 1-OH), 1.79(s, 3 H, Me19), 1.83(ddd, J = 14.5, 10.3, 2.0 Hz, 1 H, H6b), 1.97(m, 1 H, 13-OH), 2.00(s, 3 H, 7-Ac), 2.07(d, J = 1.3 Hz, 3 H, Me18), 2.26(m, 2 H, H14a, H14b), 2.29(s, 3 H, 4-Ac), 2.57(ddd, J = 14.5, 9.5, 7.3 Hz, 1 H, H6a), 4.06(d, J = 7.0 Hz, 1 H, H3), 4.17(d, J = 8.2 Hz, 1 H,

H20b), 4.31(d, J = 8.2 Hz, 1 H, H20a), 4.84(m, 1 H, H13), 4.94(dd, J = 9.5, 2.0 Hz, 1 H, H5), 5.29(s, 1 H, H10), 5.46(dd, J = 10.3, 7.3 Hz, 1 H, H7), 5.65(d, J = 7.0 Hz, 1 H, H2), 7.47(m, 2 H, benzoate, m), 7.60(m, 1 H, benzoate, p), 8.11(d, J = 8.0 Hz, 2 H, benzoate, o) ppm.

7-Troc-10-TES-10-DAB. To a mixture of 10-TES-10-DAB (40 mg, 0.061 mmol) and DMAP (72 mg, 0.61 mmol, 10 equiv) in dichloromethane (2 mL) was added trichloroethyl chloroformate (24 mL, 0.184 mmol, 3 equiv) under N₂. The reaction mixture was stirred at room temperature and the progress of the reaction was monitored by TLC analysis. After 0.5 h, TLC analysis indicated almost complete disappearance of 10-TES-10-DAB and the formation of the product as the only major spot. Methanol (5 mL) was added, and the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash chromatography using EtOAc: CH₂Cl₂ (1:10) as the eluent and dried in vacuo overnight to give 7-Troc-10-TES- 10-DAB as a white solid: yield, 49 mg (97%); ¹H NMR (500 MHz, CDCl₃) δ 0.61(m, 6 H, (CH₃CH₂)₃Si), 0.99(t, J = 7.9, 9 H, (CH₃CH₂)₃Si), 1.08(s, 3 H, Me17), 1.20(s, 3 H, Me16), 1.56(s, 1 H, 1-OH), 1.84(s, 3 H, Me19), 1.96(d, J = 4.9 Hz, 1 H, 13-OH), 2.01(ddd, J = 14.4, 10.5, 2.0 Hz, 1 H, H6b), 2.08(d, J = 1.2, 3 H, Me18), 2.29(m, 2 H, H14a, H14b), 2.29(s, 3 H, 4-Ac), 2.68(ddd, J = 14.4, 9.5, 7.3 Hz, 1 H, H6a), 4.08(d, J = 6.7 Hz, 1 H, H3), 4.18(d, J = 8.5 Hz, 1 H, H20b), 4.32(d, J = 8.5 Hz, 1 H, H20a), 4.43(d, J = 11.9 Hz, 1 H, CHH'OC(O)), 4.86(m, 1 H, H13), 4.95(dd, J = 9.3, 2.0 Hz, 1 H, H5), 4.98(d, J = 11.9 Hz, 1 H, CHH'OC(O)), 5.33(s, 1 H, H10), 5.37(dd, J = 10.5, 7.3 Hz, 1 H, H7), 5.67(d, J =

6.7 Hz, 1 H, H2), 7.48(dd, J = 7.9, 7.3 Hz, 2 H, benzoate, m), 7.60(tt, J = 7.3, 1.2 Hz, 1 H, benzoate, p), 8.11(dd, J = 7.9, 1.2 Hz, 2 H, benzoate, o)ppm.

7-*p*-Nitrobenzyloxycarbonyl-10-TES-10-DAB. To a mixture of 10-TES-10-DAB (40 mg, 0.061 mmol) and DMAP (72 mg, 0.61 mmol, 10 equiv) in dry chlorform (2 mL) was added *p*-nitrobenzyl chloroformate (131 mg, 0.61 mmol, 10 equiv) under N₂. The reaction mixture was stirred at room temperature and the progress of the reaction was monitored by TLC analysis. After 45 min, TLC analysis indicated almost complete disappearance of 10-TES-10-DAB and the formation of the product as the only major spot. Methanol (10 mL) was added, and the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash chromatography using EtOAc: CH₂Cl₂ (1:10) as the eluent and dried in vacuo overnight to give 7-*p*-Nitrobenzyloxycarbonyl-10-TES-10-DAB as a white solid: yield, 48.3 mg (95%); ¹H NMR (500 MHz, CDCl₃) δ 0.60(m, 6 H, (CH₃CH₂)₃Si), 0.95(t, J = 7.9, 9 H, (CH₃CH₂)₃Si), 1.08(s, 3 H, Me17), 1.19(s, 3 H, Me16), 1.55(s, 1 H, 1-OH), 1.83(s, 3 H, Me19), 1.93(ddd, J = 14.3, 10.4, 2.2 Hz, 1 H, H6b), 1.96(d, J = 4.9 Hz, 1 H, 13-OH), 2.09(d, J = 1.2, 3 H, Me18), 2.29(m, 2 H, H14a, H14b), 2.29(s, 3 H, 4-Ac), 2.65(ddd, J = 14.3, 9.3, 7.3 Hz, 1 H, H6a), 4.08(d, J = 7.0 Hz, 1 H, H3), 4.18(d, J = 8.6 Hz, 1 H, H20b), 4.31(d, J = 8.6 Hz, 1 H, H20a), 4.86(m, 1 H, H13), 4.95(dd, J = 9.3, 2.2 Hz, 1 H, H5), 5.06(d, J = 13.4 Hz, 1 H, CHH'OC(O)), 5.31(d, J = 13.4 Hz, 1 H, CHH'OC(O)), 5.33(s, 1 H, H10), 5.36(dd, J = 10.4, 7.3 Hz, 1 H, H7), 5.66(d, J = 7.0 Hz, 1 H, H2), 7.48(dd, J = 7.4, 7.3 Hz, 2 H, benzoate, m), 7.53(d, J =

[illegible]

7-Cbz-10-TES-10-DAB. To a mixture of 10-TES-10-DAB (40 mg, 0.061 mmol) and DMAP (440 mg, 3.64 mmol, 60 equiv) in dry chloroform (2 mL) was slowly added four equal aliquot of benzyl chloroformate (4x130 mL, 3.64 mmol, 60 equiv) via a syringe in a 10-min interval under N₂ during a period of 40 min. The reaction mixture was then stirred at room temperature and the progress of the reaction was monitored by TLC analysis. After 2 h, TLC analysis indicated almost complete disappearance of 10-TES-10-DAB and the formation of the product as the only major spot. Methanol (10 mL) was added, and the solution was poured into ethyl acetate (100 mL), washed with a saturated aqueous NaHCO₃ solution, H₂O, and brine. The solution was dried over Na₂SO₄, concentrated under reduced pressure. The residue was purified by flash chromatography using EtOAc: CH₂Cl₂ (1:10) as the eluent and dried in vacuo overnight to give 7-CBz-10-TES-10-DAB as a white solid: yield, 45 mg (93%); ¹H NMR (500 MHz, CDCl₃) δ 0.62(m, 6 H, (CH₃CH₂)₃Si), 0.97(t, J = 7.9, 9 H, (CH₃CH₂)₃Si), 1.07(s, 3 H, Me17), 1.20(s, 3 H, Me16), 1.55(s, 1 H, 1-OH), 1.81(s, 3 H, Me19), 1.91(ddd, J = 14.3, 10.5, 2.1 Hz, 1 H, H6b), 1.96(d, J = 4.9 Hz, 1 H, 13-OH), 2.10(d, J = 1.2, 3 H, Me18), 2.28(m, 2 H, H14a, H14b), 2.28(s, 3 H, 4-Ac), 2.64(ddd, J = 14.3, 9.5, 7.3 Hz, 1 H, H6a), 4.08(d, J = 7.0 Hz, 1 H, H3), 4.17(d, J = 8.5 Hz, 1 H, H20b), 4.30(d, J = 8.5 Hz, 1 H, H20a), 4.86(m, 1 H, H13), 4.95(dd, J = 9.5, 2.1 Hz, 1 H, H5), 5.01(d, J = 12.2 Hz, 1 H, CHH'OC(O)), 5.24(d, J = 12.2 Hz, 1 H, CHH'OC(O)), 5.34(s, 1 H, H10), 5.37(dd, J = 10.5, 7.3 Hz, 1 H, H7), 5.65(d, J = 7.0 Hz, 1 H, H2),

7.32-7.37 (m, 5 H, PhCH_2O), 7.47 (dd, $J = 8.3$, 7.3 Hz, 2 H, benzoate, m), 7.59 (tt, $J = 7.3$, 1.3 Hz, 1 H, benzoate, p), 8.12 (dd, $J = 8.3$, 1.3 Hz, 2 H, benzoate, o) ppm.

Example 7

Selective silylation of 10-acyl-10-DAB

7-dimethylisopropylsilyl baccatin III. To a stirred solution of baccatin III (30 mg, 0.051 mmol) in pyridine (0.6 mL) at 0 °C under N_2 , was added chlorodimethylisopropylsilane (160 μL , 1.02 mmol). The reaction mixture was stirred at that temperature and the progress of the reaction was monitored by TLC. After about 1.5 h the reaction was complete. Ethyl acetate (5 mL) was added and the solution was transferred to a separatory funnel containing 50 mL of a 50% EtOAc /Hexanes. The mixture was washed with saturated sodium bicarbonate and the organic layer was separated. The aqueous layer was extracted with 10 mL of 50% EtOAc/Hexanes and the combined organic layers were washed with saturated sodium chloride, dried over MgSO_4 , concentrated under reduced pressure. The crude product was passed through a short silica gel column to give 33.9 mg (97%) of a white solid m.p. 204-207 °C; $[\alpha]^{25}_{\text{D}} -58.6^\circ$ c (0.009, CHCl_3). ^1H NMR (CDCl_3 , 500MHz), δ 8.10 (d, $J=8.4\text{Hz}$, 2H, o-benzoate), 7.60-7.20 (m, 3H, benzoate), 6.4 (s, 1H, H10), 5.64 (d, $J=7.1\text{Hz}$, 1H, H2b), 4.95 (d, $J=4.9\text{Hz}$, 1H, H5), 4.84 (m, 1H, H13), 4.44 (dd, $J=10.4, 6.8\text{Hz}$, 1H, H7), 4.30 (d, $J=8.3\text{Hz}$, 1H, H20a), 4.14 (d, $J=8.3\text{Hz}$, 1H, H20b), 4.15 (d, $J=7.2\text{Hz}$, 1H, H3), 2.49 (m, 1H, H6a), 2.23 (m, 2H, H14's), 2.28 (s, 3H, 4Ac), 2.18 (br s, 3H, Me 18), 2.17 (s, 3H, 10Ac), 2.01 (d, $J=5.0\text{Hz}$, 13 OH), 1.86 (m, 1H, 6b), 1.69 (s, 3H, Me19), 1.61 (s, 1H, 10H), 1.20 (s, 3H, Me16), 1.05 (s, 3H, Me17), 0.87 (d, $J=7.1\text{Hz}$, 6H, i-pr), 0.73 (m, 1H, i-pr), 0.09 (s, 6H, Me2Si).

7-dimethylphenylsilyl baccatin III. To a stirred solution of baccatin III (20 mg, 0.034 mmol) in THF (1.25 mL) at -10 °C under N₂, was added chlorodimethylphenylsilane (68 uL, 0.41 mmol), followed by addition of pyridine (250 mL, 3.1 mmol). The reaction mixture was stirred at that temperature and the progress of the reaction was monitored by TLC. After about one hour the reaction was complete. Ethyl acetate (5 mL) was added and the solution was transferred to a separatory funnel containing 30 mL of 50% EtOAc /Hexanes. The mixture was washed with saturated sodium bicarbonate and the organic layer was separated. The aqueous layer was extracted with 10 mL of 50% EtOAc/Hexanes and the combined organic layers were washed with saturated sodium chloride, dried over MgSO₄, concentrated under reduced pressure. The crude product was passed through a short silica gel column to give 24.1 mg (98%) of a white solid m.p. 210-213° C; $[\alpha]_D^{25}$ -58.3.5° c (0.005, CHCl₃) ¹H NMR (CDCl₃, 500 MHz) δ 8.35(d, J=8.5 Hz, 2H, o-benzoate), 7.627.25(m, 8H, benzoate, phenyl), 6.42 (s, 1H, H10), 5.64 (d, J=6.9 Hz, 1H, H2b), 4.84(m, 1H, H5), 4.81(m, 1H, H13), 4.46 (dd, J=10.6, 6.9 Hz, 1H, H7), 4.21(d, J=8.5 Hz, 1H, H20a), 4.14 (d, J=8.5 Hz, 1H, H20b), 3.85(d, J=6.9 Hz, 1H, H3), 2.34(m, 1H, H6a), 2.26(d, J=8 Hz, 2H, H14's), 2.24(s, 3H, 4Ac), 2.15(s, 3H, 10Ac), 2.02(br d, J=1 Hz, 3H, Me 18), 1.93(d, J=5 Hz, 1H, 13OH), 1.77(m, 1H, 6b), 1.72(s, 3H, Me19), 1.59(s, 1H, 10H), 1.20(s, 3H, Me16), 1.05(s, 3H, Me17), 0.446(s, 3H, Me Si), 0.335(s, 3H, Me Si).

7-dimethylphenylsilyl-10-propionyl-10-DAB. To a stirred solution of 10-propionyl-10-DAB (0.200 g, 0.333 mmol) in THF (12 mL) at -10°C, was added chlorodimethylphenylsilane (0.668 mL, 4.00 mmol) followed by pyridine dropwise (2.48 mL, 30.64 mmol). The reaction was stirred

for 90 minutes. Ethyl acetate (20 mL) was added and the solution transferred to a separatory funnel containing 100 mL of 50% EtOAc/Hexanes. The mixture was washed with saturated sodium bicarbonate and the organic layer separated. The aqueous layer was extracted with 50%EtOAc/Hexanes (30 mL) and the combined organic extracts washed with saturated sodium chloride, dried over Na₂SO₄, concentrated *in vacuo*. The crude solid was then purified with flash column chromatography using 50% EtOAc/hexane as eluent to give 7-dimethylphenylsilyl-10-propionyl-10-DAB (0.242 g, 99%) as a solid. ¹H NMR (CDCl₃, 500 MHz), δ 0.34, 0.45(2 s, 6 H, Me₂Si), 1.05(s, 3 H, Me17), 1.20(t, J = 7.5 Hz, 3 H, CH₃CH₂), 1.21(s, 3 H, Me16), 1.60(s, 1 H, 1-OH), 1.72(s, 3 H, Me19), 1.78(ddd, J = 14.5, 10.0, 2.0 Hz, 1 H, H6b), 2.04(m, 1 H, 13-OH), 2.05(s, 3 H, Me18), 2.27(m, 2 H, H14a, H14b), 2.25(s, 3 H, 4-Ac), 2.34(ddd, J = 14.5, 9.5, 7.0 Hz, 1 H, H6a), 2.42, 2.49(2 dq, J = 16.5, 7.5 Hz, 6 H, CH₃CH₂), 3.87(d, J = 7.5 Hz, 1 H, H3), 4.14(d, J = 8.0 Hz, 1 H, H20b), 4.27(d, J = 8.0 Hz, 1 H, H20a), 4.47(dd, J = 10.0, 7.0 Hz, 1 H, H7), 4.82(m, 1 H, H13), 4.85(dd, J = 9.5, 2.0 Hz, 1 H, H5), 5.64(d, J = 7.5 Hz, 1 H, H2), 6.44(s, 1 H, H10), 7.32-7.36, 7.55-7.57(2 m, 5 H, PhSi), 7.46(m, 2 H, benzoate, m), 7.59(m, 1 H, benzoate, p), 8.10(d, J = 8.0 Hz, 2 H, benzoate, o) ppm.

7-Dimethylphenylsilyl-10-cyclopropanecarbonyl-10-DAB. To a solution of 10-cyclopropanecarbonyl-10-DAB (680 mg, 1.1 mmol) in THF (25 mL) were added with stirring pyridine (3.5 mL) and then chlorodimethylphenylsilane (1.8 mL, 11 mmol) at -10 °C under N₂. The solution was stirred till the reaction completed. Then quenched with sat. NaHCO₃ (20 mL). The mixture was extracted with EtOAc (2 x 250 mL). The combined organic

layers were washed with brine (2 x 10 mL), dried and filtered. Concentration of the filtrate in *vacuo* and followed by flash chromatography (hexane:EtOAc, 4:1) gave 7-Dimethyl-phenylsilyl-10-cyclopropane-carbonyl-10-DAB (816 mg, ~100%). ¹H NMR (CDCl₃, 500 MHz), δ 0.32, 0.43 (2 s, 6 H, Me₂Si), 0.91, 1.00, 1.17 (3 m, 5 H, cyclopropyl), 1.07 (s, 3 H, Me17), 1.21 (s, 3 H, Me16), 1.73 (s, 3 H, Me19), 1.74 (s, 1 H, 1-OH), 1.78 (ddd, J = 14.4, 10.5, 2.1 Hz, 1 H, H6b), 2.04 (m, 1 H, 13-OH), 2.05 (d, J = 1.5 Hz, 3 H, Me18), 2.24 (s, 3 H, 4-Ac), 2.26 (m, 2 H, H14a, H14b), 2.34 (ddd, J = 14.4, 9.5, 6.7 Hz, 1 H, H6a), 3.87 (d, J = 7.0 Hz, 1 H, H3), 4.15 (d, J = 8.2 Hz, 1 H, H20b), 4.26 (d, J = 8.2 Hz, 1 H, H20a), 4.46 (dd, J = 10.5, 6.7 Hz, 1 H, H7), 4.82 (m, 1 H, H13), 4.85 (dd, J = 9.5, 2.1 Hz, 1 H, H5), 5.65 (d, J = 7.0 Hz, 1 H, H2), 6.44 (s, 1 H, H10), 7.32-7.36, 7.55-7.57 (2 m, 5 H, PhSi), 7.46 (m, 2 H, benzoate, m), 7.59 (m, 1 H, benzoate, p), 8.10 (d, J = 8.0 Hz, 2 H, benzoate, o) ppm.

Example 8

7-*p*-Nitrobenzyloxycarbonyl-10-DAB. To a THF solution (1 mL) of 10-alloc-7-*p*-nitrobenzyloxycarbonyl-10-DAB (34 mg, 0.048 mmol) at room temperature was added a THF solution (1 mL) of formic acid (19 mL, 0.48 mmol, 10 equiv) and butylamine (47 mL, 0.48 mmol, 10 equiv), followed by the addition of Pd(PPh₃)₄ under N₂. The reaction mixture was stirred at room temperature for 0.5 h. EtOAc (10 mL) was added, and the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:2) as the eluent and dried in *vacuo* to give 7-*p*-nitrobenzyloxycarbonyl-10-DAB as a colorless

solid: yield, 28 mg (93%). $[\alpha]_{\text{Hg}} -38^\circ$ (CHCl_3 , $c = 0.48$); ^1H NMR (400 MHz, CDCl_3) δ 1.06(s, 3 H, Me16), 1.09(s, 3 H, Me17), 1.55(s, 1 H, 1-OH), 1.86(s, 3 H, Me19), 2.01(ddd, $J = 14.4, 10.7, 2.0$ Hz, 1 H, H6b), 2.03(d, $J = 5.1$ Hz, 1 H, 13-OH), 2.09(d, $J = 1.3, 3$ H, Me18), 2.28(m, 2 H, H14a, H14b), 2.30(s, 3 H, 4-Ac), 2.62(ddd, $J = 14.4, 9.5, 7.3$ Hz, 1 H, H6a), 3.89(d, $J = 2.0$ Hz, 1 H, 10-OH), 4.08(d, $J = 6.9$ Hz, 1 H, H3), 4.20(d, $J = 8.4$ Hz, 1 H, H20b), 4.34(d, $J = 8.4$ Hz, 1 H, H20a), 4.88(m, 1 H, H13), 4.96(dd, $J = 9.5, 2.0$ Hz, 1 H, H5), 5.19(d, $J = 13.3, 1$ H, $\text{CHH}'\text{OC}(\text{O})$), 5.26(d, $J = 13.3, 1$ H, $\text{CHH}'\text{OC}(\text{O})$), 5.36(dd, $J = 10.7, 7.3$ Hz, 1 H, H7), 5.40(d, $J = 2.0$ Hz, 1 H, H10), 5.64(d, $J = 6.9$ Hz, 1 H, H2), 7.48(dd, $J = 8.1, 7.5$ Hz, 2 H, benzoate, m), 7.52(d, $J = 8.7, 2$ H, $\text{NO}_2\text{C}_6\text{H}_4$), 7.61(tt, $J = 7.5, 1.3$ Hz, 1 H, benzoate, p), 8.10(d, $J = 8.1, 1.3$ Hz, 2 H, benzoate, o), 8.26(d, $J = 8.7, 2$ H, $\text{NO}_2\text{C}_6\text{H}_4$) ppm. ^{13}C NMR (75 MHz, CDCl_3) δ 10.5(Me(19)), 14.6(Me(18)), 19.4(4-Ac), 22.2, 26.4(Me16, Me17), 33.2(C(6)), 38.7(C(14)), 42.4(C(15)), 46.5(C(3)), 56.5(C(8)), 67.9, 68.3(C(13), $\text{OCH}_2\text{Ph-NO}_2$ -p), 74.7, 75.2, 76.8(C(7), C(2), C(10), C(20)), 78.8(C(1)), 80.4(C(4)), 83.6(C(5)), 124.1, 128.4, 128.9, 130.3, 133.9 ($\text{OCH}_2\text{Ph-NO}_2$ -p, benzoate), 135.0(C(11)), 142.4, 143.0 ($\text{OCH}_2\text{Ph-NO}_2$ -p, C(12)), 154.2(OC(O)O), 167.3(benzoate), 171.1(4-Ac), 211.6(C(9)) ppm.

Example 9

Selective Esterification of the C-10 Hydroxyl of 10-DAB using the catalytic DyCl_3 reaction: A solution of butyric anhydride (0.55 mmol) in THF (1.32 ml) was added, under a nitrogen atmosphere, to a solid mixture of 10-DAB (30 mg, 0.055 mmol) and DyCl_3 (1.3 mg, 10 mol.% wrt 10-DAB). The resulting suspension was stirred at room temperature until judged complete by TLC (2:1

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EtOAc/Hexane). The reaction was diluted with EtOAc and washed three times with saturated NaHCO_3 solution. The combined bicarbonate washings were extracted three times with EtOAc, these combined organics were dried (Na_2SO_4) and the solvent evaporated. The crude product was triturated with hexanes and the mother liquors decanted away. Crystallization from EtOAc/hexanes yielded 10-butyl-10-DAB identical to that isolated from the CeCl_3 catalysed reaction.

Example 10

Selective Esterification of the C-10 Hydroxyl of 10-DAB using the catalytic YbCl_3 reaction: A solution of butyric anhydride (0.55 mmol) in THF (1.32 ml) was added, under a nitrogen atmosphere, to a solid mixture of 10-DAB (30 mg, 0.055 mmol) and YbCl_3 (1.3 mg, 10 mol.% wrt 10-DAB). The resulting suspension was stirred at room temperature until judged complete by TLC (2:1 EtOAc/Hexane). The reaction was diluted with EtOAc and washed three times with saturated NaHCO_3 solution. The combined bicarbonate washings were extracted three times with EtOAc, these combined organics were dried (Na_2SO_4) and the solvent evaporated. The crude product was triturated with hexanes and the mother liquors decanted away. Crystallization from EtOAc/hexanes yielded 10-butyl-10-DAB identical to that isolated from the CeCl_3 catalysed reaction.

In view of the above, it will be seen that the several objects of the invention are achieved.

As various changes could be made in the above compositions without departing from the scope of the invention, it is intended that all matter contained in

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1. **Introduction**
 2. **Background**
 3. **Methodology**
 4. **Results**
 5. **Discussion**
 6. **Conclusion**
 7. **References**
 8. **Appendix**
 9. **Figure 1**
 10. **Figure 2**
 11. **Figure 3**
 12. **Figure 4**
 13. **Figure 5**
 14. **Figure 6**
 15. **Figure 7**
 16. **Figure 8**
 17. **Figure 9**
 18. **Figure 10**
 19. **Figure 11**
 20. **Figure 12**
 21. **Figure 13**
 22. **Figure 14**
 23. **Figure 15**
 24. **Figure 16**
 25. **Figure 17**
 26. **Figure 18**
 27. **Figure 19**
 28. **Figure 20**
 29. **Figure 21**
 30. **Figure 22**
 31. **Figure 23**
 32. **Figure 24**
 33. **Figure 25**
 34. **Figure 26**
 35. **Figure 27**
 36. **Figure 28**
 37. **Figure 29**
 38. **Figure 30**
 39. **Figure 31**
 40. **Figure 32**
 41. **Figure 33**
 42. **Figure 34**
 43. **Figure 35**
 44. **Figure 36**
 45. **Figure 37**
 46. **Figure 38**
 47. **Figure 39**
 48. **Figure 40**
 49. **Figure 41**
 50. **Figure 42**
 51. **Figure 43**
 52. **Figure 44**
 53. **Figure 45**
 54. **Figure 46**
 55. **Figure 47**
 56. **Figure 48**
 57. **Figure 49**
 58. **Figure 50**
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 60. **Figure 52**
 61. **Figure 53**
 62. **Figure 54**
 63. **Figure 55**
 64. **Figure 56**
 65. **Figure 57**
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 67. **Figure 59**
 68. **Figure 60**
 69. **Figure 61**
 70. **Figure 62**
 71. **Figure 63**
 72. **Figure 64**
 73. **Figure 65**
 74. **Figure 66**
 75. **Figure 67**
 76. **Figure 68**
 77. **Figure 69**
 78. **Figure 70**
 79. **Figure 71**
 80. **Figure 72**
 81. **Figure 73**
 82. **Figure 74**
 83. **Figure 75**
 84. **Figure 76**
 85. **Figure 77**
 86. **Figure 78**
 87. **Figure 79**
 88. **Figure 80**
 89. **Figure 81**
 90. **Figure 82**
 91. **Figure 83**
 92. **Figure 84**
 93. **Figure 85**
 94. **Figure 86**
 95. **Figure 87**
 96. **Figure 88**
 97. **Figure 89**
 98. **Figure 90**
 99. **Figure 91**
 100. **Figure 92**
 101. **Figure 93**
 102. **Figure 94**
 103. **Figure 95**
 104. **Figure 96**
 105. **Figure 97**
 106. **Figure 98**
 107. **Figure 99**
 108. **Figure 100**
 109. **Figure 101**
 110. **Figure 102**
 111. **Figure 103**
 112. **Figure 104**
 113. **Figure 105**
 114. **Figure 106**
 115. **Figure 107**
 116. **Figure 108**
 117. **Figure 109**
 118. **Figure 110**
 119. **Figure 111**
 120. **Figure 112**
 121. **Figure 113**
 122. **Figure 114**
 123. **Figure 115**
 124. **Figure 116**
 125. **Figure 117**
 126. **Figure 118**
 127. **Figure 119**
 128. **Figure 120**
 129. **Figure 121**
 130. **Figure 122**
 131. **Figure 123**
 132. **Figure 124**
 133. **Figure 125**
 134. **Figure 126**
 135. **Figure 127**
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 140. **Figure 132**
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 147. **Figure 139**
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 149. **Figure 141**
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 151. **Figure 143**
 152. **Figure 144**
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 156. **Figure 148**
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 159. **Figure 151**
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 161. **Figure 153**
 162. **Figure 154**
 163. **Figure 155**
 164. **Figure 156**
 165. **Figure 157**
 166. **Figure 158**
 167. **Figure 159**
 168. **Figure 160**
 169. **Figure 161**
 170. **Figure 162**
 171. **Figure 163**
 172. **Figure 164**
 173. **Figure 165**
 174. **Figure 166**
 175. **Figure 167**
 176. **Figure 168**
 177. **Figure 169**
 178. **Figure 170**
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 181. **Figure 173**
 182. **Figure 174**
 183. **Figure 175**
 184. **Figure 176**
 185. **Figure 177**
 186. **Figure 178**
 187. **Figure 179**
 188. **Figure 180**
 189. **Figure 181**
 190. **Figure 182**
 191. **Figure 183**
 192. **Figure 184**
 193. **Figure 185**
 194. **Figure 186**
 195. **Figure 187**
 196. **Figure 188**
 197. **Figure 189**
 198. **Figure 190**
 199. **Figure 191**
 200. **Figure 192**
 201. **Figure 193**
 202. **Figure 194**
 203. **Figure 195**
 204. **Figure 196**
 205. **Figure 197**
 206. **Figure 198**
 207. **Figure 199**
 208. **Figure 200**
 209. **Figure 201**
 210. **Figure 202**
 211. **Figure 203**
 212. **Figure 204**
 213. **Figure 205**
 214. **Figure 206**
 215. **Figure 207**
 216. **Figure 208**
 217. **Figure 209**

What is claimed is:

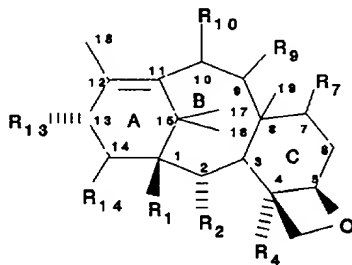
1. A process for the acylation of a C(10) hydroxy group of a taxane, the process comprising

treating the taxane with an acylating agent in a reaction mixture containing less than one equivalent of an amine base for each equivalent of taxane to form a C(10) acylated taxane.

2. The process of claim 1 wherein the taxane has C(7) and C(10) hydroxy groups and the acylating agent selectively reacts with the C(10) hydroxy group.

3. The process of claim 1 wherein the taxane reacted with the acylating agent is 10-deacetyl baccatin III.

4. The process of claim 1 wherein the taxane has the structure:



wherein

5 R_1 is hydrogen, hydroxy, protected hydroxy, or together with R_{14} or R_2 forms a carbonate;

R_2 is keto, $-OT_2$, acyloxy, or together with R_1 forms a carbonate;

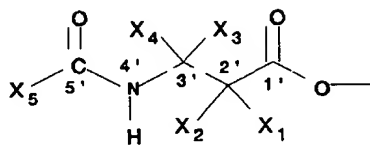
R_4 is $-OT_4$ or acyloxy;

10 R_7 is hydrogen, halogen, $-OT_7$, $-OCOZ_7$, or $-OCOOZ_7$;

R_9 is hydrogen, keto, $-OT_9$, $-OCOZ_9$, or $-OCOOZ_9$;

R_{10} is hydroxy;

R_{13} is hydroxy, protected hydroxy, keto, or



- 15 R_{14} is hydrogen, $-OT_{14}$, acyloxy, or together with R_1 forms a carbonate;
 T_2 , T_4 , T_7 , T_9 and T_{14} are independently hydrogen or hydroxy protecting group;
 X_1 is $-OX_6$, $-SX_7$, or $-NX_8X_9$;
20 X_2 is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;
 X_3 and X_4 are independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;
 X_5 is $-X_{10}$, $-OX_{10}$, $-SX_{10}$, $-NX_8X_{10}$, or $-SO_2X_{11}$;
25 X_6 is hydrocarbyl, substituted hydrocarbyl, heteroaryl, hydroxy protecting group or a functional group which increases the water solubility of the taxane derivative;
 X_7 is hydrocarbyl, substituted hydrocarbyl, heteroaryl, or sulfhydryl protecting group;
30 X_8 is hydrogen, hydrocarbyl, or substituted hydrocarbyl;
 X_9 is an amino protecting group;
 X_{10} is hydrocarbyl, substituted hydrocarbyl, or heteroaryl;
35 X_{11} is hydrocarbyl, substituted hydrocarbyl, heteroaryl, $-OX_{10}$, or $-NX_8X_{14}$; and
 X_{14} is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

5. The process of claim 4 wherein

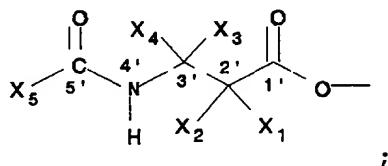
R_1 is hydroxy or together with R_{14} or R_2 forms a carbonate;

R_2 is $-\text{OCOZ}_2$, $-\text{OCOOZ}_2$, or together with R_1 forms a
5 carbonate;

R_4 is $-\text{OCOZ}_4$;

R_9 is hydrogen or keto;

R_{13} is hydroxy, protected hydroxy, or



10 R_{14} is hydrogen, hydroxy, protected hydroxy, or
together with R_1 forms a carbonate;

X_1 is $-\text{OX}_6$ or $-\text{NX}_8\text{X}_9$;

X_2 is hydrogen, hydrocarbyl, or substituted
hydrocarbyl;

15 X_3 and X_4 are independently hydrogen, hydrocarbyl,
substituted hydrocarbyl, or heteroaryl;

X_5 is $-\text{X}_{10}$, $-\text{OX}_{10}$, or $-\text{NX}_8\text{X}_{10}$;

X_6 is a hydroxy protecting group;

20 X_8 is hydrogen, hydrocarbyl, or substituted
hydrocarbyl;

X_9 is an amino protecting group;

X_{10} is hydrocarbyl, substituted hydrocarbyl, or
heteroaryl; and

25 Z_2 and Z_4 are independently hydrocarbyl, substituted
hydrocarbyl, or heteroaryl.

6. The process of claim 1 wherein the acylating
agent is selected from the group consisting of

anhydrides, dicarbonates, thiodicarbonates, and isocyanates.

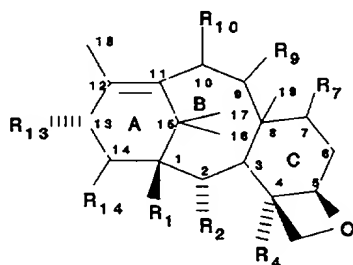
7. The process of claim 6 wherein the reaction mixture contains a Lewis acid.

8. The process of claim 7 wherein the taxane reacted with the acylating agent is 10-deacetyl baccatin III.

9. The process of claim 1 wherein the reaction mixture contains a Lewis acid.

10. The process of claim 9 wherein the taxane reacted with the acylating agent is 10-deacetyl baccatin III.

11. The process of claim 9 wherein the taxane has the structure:



wherein

5 R_1 is hydrogen, hydroxy, protected hydroxy, or together with R_{14} or R_2 forms a carbonate;

R_2 is keto, $-OT_2$, acyloxy, or together with R_1 forms a carbonate;

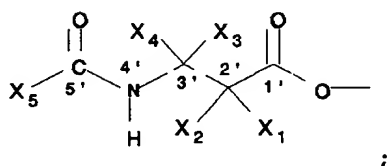
R_4 is $-OT_4$ or acyloxy;

10 R_7 is hydrogen, halogen, $-OT_7$, $-OCOZ_7$, or $-OCOOZ_7$;

R_9 is hydrogen, keto, $-OT_9$, $-OCOZ_9$, or $-OCOOZ_9$;

R_{10} is hydroxy;

R_{13} is hydroxy, protected hydroxy, keto, or



15 R_{14} is hydrogen, $-OT_{14}$, acyloxy, or together with R_1 forms a carbonate;

T_2 , T_4 , T_7 , T_9 and T_{14} are independently hydrogen or hydroxy protecting group;

X_1 is $-OX_6$, $-SX_7$, or $-NX_8X_9$;

20 X_2 is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

X_3 and X_4 are independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

X_5 is $-X_{10}$, $-OX_{10}$, $-SX_{10}$, $-NX_8X_{10}$, or $-SO_2X_{11}$;

25 X_6 is hydrocarbyl, substituted hydrocarbyl, heteroaryl, hydroxy protecting group or a functional group which increases the water solubility of the taxane derivative;

30 X_7 is hydrocarbyl, substituted hydrocarbyl, heteroaryl, or sulfhydryl protecting group;

X_8 is hydrogen, hydrocarbyl, or substituted hydrocarbyl;

X_9 is an amino protecting group;

35 X_{10} is hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

X_{11} is hydrocarbyl, substituted hydrocarbyl, heteroaryl, $-OX_{10}$, or $-NX_8X_{14}$; and

X_{14} is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

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12. The process of claim 11 wherein

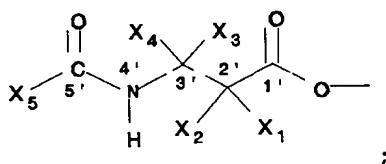
R_1 is hydroxy or together with R_{14} or R_2 forms a carbonate;

R_2 is $-\text{OCOZ}_2$, $-\text{OCOOZ}_2$, or together with R_1 forms a
5 carbonate;

R_4 is $-\text{OCOZ}_4$;

R_9 is hydrogen or keto;

R_{13} is hydroxy, protected hydroxy, or



10 R_{14} is hydrogen, hydroxy, protected hydroxy, or
together with R_1 forms a carbonate;

X_1 is $-\text{OX}_6$ or $-\text{NX}_8\text{X}_9$;

X_2 is hydrogen, hydrocarbyl, or substituted
hydrocarbyl;

15 X_3 and X_4 are independently hydrogen, hydrocarbyl,
substituted hydrocarbyl, or heteroaryl;

X_5 is $-\text{X}_{10}$, $-\text{OX}_{10}$, or $-\text{NX}_8\text{X}_{10}$;

X_6 is a hydroxy protecting group;

20 X_8 is hydrogen, hydrocarbyl, or substituted
hydrocarbyl;

X_9 is an amino protecting group;

X_{10} is hydrocarbyl, substituted hydrocarbyl, or
heteroaryl; and

25 Z_2 and Z_4 are independently hydrocarbyl, substituted
hydrocarbyl, or heteroaryl.

13. The process of claim 9 wherein the Lewis acid
is selected from the group consisting of the halides or

triflates of the Group IB, IIB, IIIB, IVB, VB, VIIB, VIII, IIIA, IVA, lanthanide and actinide elements.

14. The process of claim 13 wherein the Lewis acid is selected from the group consisting of zinc chloride, stannic chloride, cerium trichloride, cuprous chloride, lanthanum trichloride, dysprosium trichloride and
5 ytterbium trichloride.

15. The process of claim 1 wherein the C(10) acylated taxane comprises a C(7) hydroxy group and the process additionally comprises treating the C(10) acylated taxane with a silylating agent to silylate the
5 C(7) hydroxy group.

16. The process of claim 15 wherein the C(10) acylated taxane is baccatin III.

17. The process of claim 1 wherein the C(10) acylated taxane comprises a C(7) hydroxy group and the process additionally comprises treating the C(10) acylated taxane with an acylating agent to acylate the
5 C(7) hydroxy group.

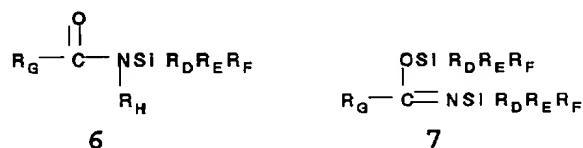
18. The process of claim 17 wherein the C(10) acylated taxane is baccatin III.

19. The process of claim 1 wherein the C(10) acylated taxane comprises a C(13) hydroxy, metallic oxide, or ammonium oxide substituent and the process additionally comprises the step of esterifying the C(10)
5 acylated taxane by treating the C(10) acylated taxane with a side chain precursor selected from the group consisting of β -lactams, oxazolines, oxazolidine

carboxylic acids, oxazolidine carboxylic acid anhydrides, and isoserine derivatives.

20. A process for the silylation of a C(10) hydroxy group of a taxane, the process comprising treating the taxane with a silylamide or a bissilylamide to form a C(10) silylated taxane.

21. The process of claim 20 wherein the silylamide or bissilylamide corresponds to structures 6 or 7, respectively:



wherein R_D , R_E , R_F , R_G , and R_H are independently hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

22. The process of claim 21 wherein the taxane is treated with the silylamide or bissilylamide in the presence of an alkali metal base catalyst.

23. The process of claim 21 wherein the taxane is treated with the silylamide or bissilylamide in the presence of a catalyst selected from the group consisting of lithium amide catalysts.

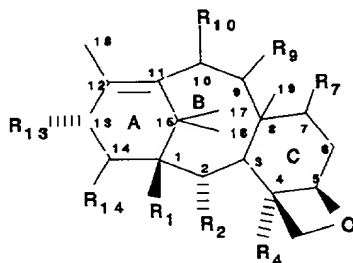
24. The process of claim 20 wherein the silylamide or bissilylamide is selected from the group consisting of tri(hydrocarbyl)silyl-trifluoromethylacetamides and bis tri(hydrocarbyl)-silyltrifluoromethylacetamides, with the

- 5 hydrocarbyl moiety being substituted or unsubstituted alkyl or aryl.

25. The process of claim 20 wherein the C(10) silylated taxane comprises a C(7) hydroxy group and the process additionally comprises treating the C(10) silylated taxane with an acylating agent to acylate the C(7) hydroxy group.

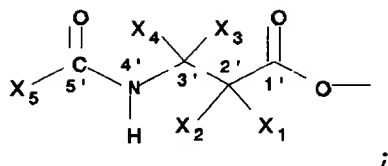
26. The process of claim 20 wherein the taxane treated with the silylamide or bissilylamide is 10-deacetyl baccatin III and the process additionally comprises treating the C(10) silylated taxane with an acylating agent to acylate the C(7) hydroxy group.

27. The process of claim 20 wherein the taxane has the structure:



wherein

- 5 R_1 is hydrogen, hydroxy, protected hydroxy, or together with R_{14} or R_2 forms a carbonate;
 R_2 is keto, $-OT_2$, acyloxy or together with R_1 forms a carbonate;
 R_4 is $-OT_4$ or acyloxy;
10 R_7 is hydrogen, halogen, $-OT_7$, or acyloxy;
 R_9 is hydrogen, keto, $-OT_9$, or acyloxy;
 R_{10} is hydroxy;
 R_{13} is hydroxy, protected hydroxy, keto, or



15 R_{14} is hydrogen, $-OT_{14}$, acyloxy, or together with R_1
forms a carbonate;
 T_2 , T_4 , T_7 , T_9 , and T_{14} are independently hydrogen or
hydroxy protecting group;
 X_1 is $-OX_6$, $-SX_7$, or $-NX_8X_9$;
20 X_2 is hydrogen, hydrocarbyl, substituted hydrocarbyl,
or heteroaryl;
 X_3 and X_4 are independently hydrogen, hydrocarbyl,
substituted hydrocarbyl, or heteroaryl;
 X_5 is $-X_{10}$, $-OX_{10}$, $-SX_{10}$, $-NX_8X_{10}$,
25 or $-SO_2X_{11}$;
 X_6 is hydrocarbyl, substituted hydrocarbyl,
heteroaryl, hydroxy protecting group or a functional
group which increases the water solubility of the taxane
derivative;
30 X_7 is hydrocarbyl, substituted hydrocarbyl,
heteroaryl, or sulfhydryl protecting group;
 X_8 is hydrogen, hydrocarbyl, or substituted
hydrocarbyl;
 X_9 is an amino protecting group;
35 X_{10} is hydrocarbyl, substituted hydrocarbyl, or
heteroaryl;
 X_{11} is hydrocarbyl, substituted hydrocarbyl,
heteroaryl, $-OX_{10}$, or $-NX_8X_{14}$; and
 X_{14} is hydrogen, hydrocarbyl, substituted
40 hydrocarbyl, or heteroaryl.

28. The process of claim 27 wherein

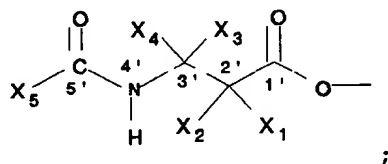
R_1 is hydroxy or together with R_{14} or R_2 forms a carbonate;

R_2 is $-\text{OCOZ}_2$, $-\text{OCOOZ}_2$, or together with R_1 forms a
5 carbonate;

R_4 is $-\text{OCOZ}_4$;

R_9 is hydrogen or keto;

R_{13} is hydroxy, protected hydroxy, or



10 R_{14} is hydrogen, hydroxy, protected hydroxy, or
together with R_1 forms a carbonate;

X_1 is $-\text{OX}_6$ or $-\text{NX}_8\text{X}_9$;

X_2 is hydrogen, hydrocarbyl, or substituted
hydrocarbyl;

15 X_3 and X_4 are independently hydrogen, hydrocarbyl,
substituted hydrocarbyl, or heteroaryl;

X_5 is $-\text{X}_{10}$, $-\text{OX}_{10}$, or $-\text{NX}_8\text{X}_{10}$;

X_6 is a hydroxy protecting group;

X_8 is hydrogen, hydrocarbyl, or substituted
20 hydrocarbyl;

X_9 is an amino protecting group;

X_{10} is hydrocarbyl, substituted hydrocarbyl, or
heteroaryl; and

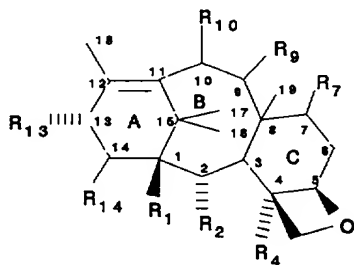
Z_2 and Z_4 are independently hydrocarbyl, substituted
25 hydrocarbyl, or heteroaryl.

29. A process for converting the C(7) hydroxy group
of a 10-acyloxy-7-hydroxytaxane to an acetal or ketal,
the process comprising treating the 10-acyloxy-7-

hydroxytaxane with a ketalizing agent in the presence of
5 an acid catalyst to form a C(10) ketalized taxane.

30. The process of claim 29 wherein the C(10)
substituent of the 10-acyloxy-7-hydroxytaxane is acetoxy.

31. The process of claim 29 wherein the taxane has
the structure:



wherein

5 R_1 is hydrogen, hydroxy, protected hydroxy, or
together with R_{14} or R_2 forms a carbonate;

R_2 is keto, $-OT_2$, acyloxy, or together with R_1 forms
a carbonate;

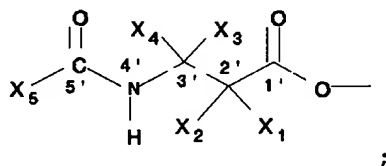
R_4 is $-OT_4$, or acyloxy;

10 R_7 is hydroxy;

R_9 is hydrogen, keto, $-OT_9$, or acyloxy;

R_{10} is acyloxy;

R_{13} is hydroxy, protected hydroxy, keto, or



15 R_{14} is hydrogen, $-OT_4$, acyloxy, or together with R_1
forms a carbonate;

T_2 , T_4 , T_9 , and T_{14} are independently hydrogen or hydroxy protecting group;

X_1 is $-OX_6$, $-SX_7$, or $-NX_8X_9$;

20 X_2 is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

X_3 and X_4 are independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

25 X_5 is $-X_{10}$, $-OX_{10}$, $-SX_{10}$, $-NX_8X_{10}$, or $-SO_2X_{11}$;

X_6 is hydrocarbyl, substituted hydrocarbyl, heteroaryl, hydroxy protecting group or a functional group which increases the water solubility of the taxane derivative;

30 X_7 is hydrocarbyl, substituted hydrocarbyl, heteroaryl, or sulfhydryl protecting group;

X_8 is hydrogen, hydrocarbyl, or substituted hydrocarbyl;

X_9 is an amino protecting group;

35 X_{10} is hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

X_{11} is hydrocarbyl, substituted hydrocarbyl, heteroaryl, $-OX_{10}$, or $-NX_8X_{14}$; and

40 X_{14} is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

32. The process of claim 31 wherein

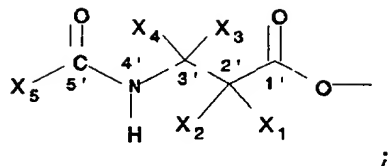
R_1 is hydroxy or together with R_{14} or R_2 forms a carbonate;

5 R_2 is $-OCOZ_2$, $-OCOOZ_2$, or together with R_1 forms a carbonate;

R_4 is $-OCOZ_4$;

R_9 is hydrogen or keto;

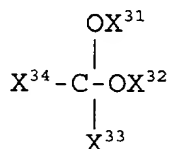
R_{13} is hydroxy, protected hydroxy, or



- 10 R₁₄ is hydrogen, hydroxy, protected hydroxy, or
together with R₁ forms a carbonate;
 X₁ is -OX₆ or -NX₈X₉;
 X₂ is hydrogen, hydrocarbyl, or substituted
hydrocarbyl;
15 X₃ and X₄ are independently hydrogen, hydrocarbyl,
substituted hydrocarbyl, or heteroaryl;
 X₅ is -X₁₀, -OX₁₀, or -NX₈X₁₀;
 X₆ is a hydroxy protecting group;
 X₈ is hydrogen, hydrocarbyl, or substituted
20 hydrocarbyl;
 X₉ is an amino protecting group;
 X₁₀ is hydrocarbyl, substituted hydrocarbyl, or
heteroaryl; and
 Z₂ and Z₄ are independently hydrocarbyl, substituted
25 hydrocarbyl, or heteroaryl.

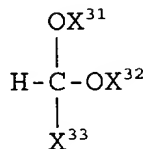
33. The process of claim 29 wherein the acid catalyst is an inorganic acid.

34. The process of claim 29 wherein the ketalizing agent has the formula



wherein X^{31} , X^{32} , X^{33} and X^{34} are independently hydrocarbyl, substituted hydrocarbyl or heteroaryl.

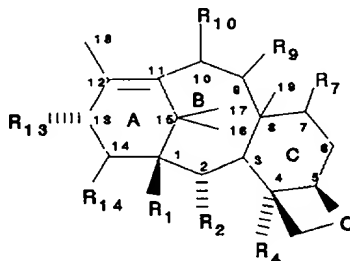
35. The process of claim 29 wherein the ketalizing agent has the formula



wherein X^{31} , X^{32} and X^{33} are independently hydrocarbyl, substituted hydrocarbyl or heteroaryl.

36. The process of claim 29 wherein the ketalizing agent is a vinyl ether.

37. A taxane having the structure:



5

wherein

M is a metal or comprises ammonium:

R_1 is hydrogen, hydroxy, protected hydroxy, or together with R_{14} or R_2 forms a carbonate;

10 R_2 is keto, $-\text{OT}_2$, acyloxy, or together with R_1 forms a carbonate;

R_4 is $-\text{OT}_4$ or acyloxy;

R_7 is $-\text{OSiR}_j\text{R}_k\text{R}_l$;

R_9 is hydrogen, keto, $-\text{OT}_9$, or acyloxy;

15 R_{10} is hydrogen, keto, $-\text{OT}_{10}$, or acyloxy;

R_{13} is hydroxy, protected hydroxy, keto, or $\text{MO}-$;

R_{14} is hydrogen, $-\text{OT}_{14}$, acyloxy, or together with R_1 forms a carbonate;

20 R_J , R_K , R_L are independently hydrocarbyl, substituted hydrocarbyl, or heteroaryl, provided, however, if each of R_J , R_K and R_L are alkyl, at least one of R_J , R_K and R_L comprises a carbon skeleton having at least four carbon atoms; and

25 T_2 , T_4 , T_9 , T_{10} , and T_{14} are independently hydrogen or hydroxy protecting group.

38. The taxane of claim 37 wherein R_{10} is $-OT_{10}$, T_{10} is $SiR_DR_ER_F$, and R_D , R_E , R_F are independently hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

39. The taxane of claim 37 wherein

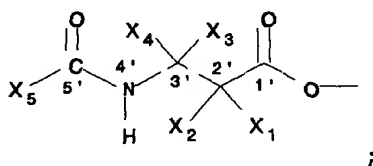
R_1 is hydroxy or together with R_{14} or R_2 forms a carbonate;

5 R_2 is $-OCOZ_2$, $-OCOOZ_2$, or together with R_1 forms a carbonate;

R_4 is $-OCOZ_4$;

R_9 is hydrogen or keto;

R_{13} is hydroxy, protected hydroxy, or



10 R_{14} is hydrogen, hydroxy, protected hydroxy, or together with R_1 forms a carbonate;

X_1 is $-OX_6$ or $-NX_8X_9$;

X_2 is hydrogen, hydrocarbyl, or substituted hydrocarbyl;

15 X_3 and X_4 are independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

X_5 is $-X_{10}$, $-OX_{10}$, or $-NX_8X_{10}$;

X_6 is a hydroxy protecting group;

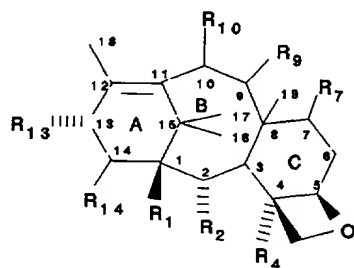
X_8 is hydrogen, hydrocarbyl, or substituted
20 hydrocarbyl;

X_9 is an amino protecting group;

X_{10} is hydrocarbyl, substituted hydrocarbyl, or
heteroaryl; and

Z_2 and Z_4 are independently hydrocarbyl, substituted
25 hydrocarbyl, or heteroaryl.

40. A taxane having the structure:



wherein

M is a metal or comprises ammonium:

5 R_1 is hydrogen, hydroxy, protected hydroxy, or
together with R_{14} or R_2 forms a carbonate;

R_2 is keto, $-OT_2$, acyloxy, or together with R_1 forms
a carbonate;

R_4 is $-OT_4$ or acyloxy;

10 R_7 is $-OT_7$ or acyloxy;

R_9 is hydrogen, keto, $-OT_9$, or acyloxy;

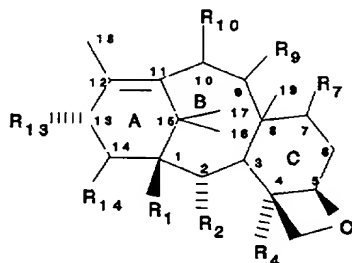
R_{10} is acyloxy, provided that the acyl moiety is
other than acetoxy;

R_{13} is hydroxy, protected hydroxy, keto, or $MO-$;

15 R_{14} is hydrogen, $-OT_{14}$, acyloxy, or together with R_1
forms a carbonate; and

T_2 , T_4 , T_7 , T_9 , and T_{14} are independently hydrogen or
hydroxy protecting group.

41. A taxane having the structure:



wherein

M is a metal or comprises ammonium:

5 R_1 is hydrogen, hydroxy, protected hydroxy, or
together with R_{14} or R_2 forms a carbonate;

R_2 is keto, $-OT_2$, acyloxy, or together with R_1 forms
a carbonate;

R_4 is $-OT_4$ or acyloxy;

10 R_7 is hydrogen, halogen, $-OT_7$, or acyloxy;

R_9 is hydrogen, keto, $-OT_9$, or acyloxy;

R_{10} is $-OSiR_D R_E R_F$;

R_{13} is hydroxy, protected hydroxy, keto, or $MO-$;

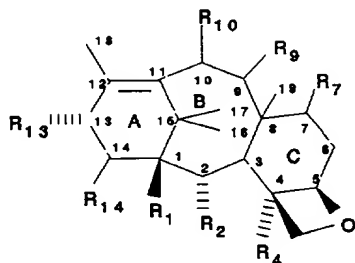
15 R_{14} is hydrogen, $-OT_{14}$ or together with R_1 forms a
carbonate;

R_D , R_E , R_F are independently hydrocarbyl, substituted
hydrocarbyl, or heteroaryl; and

T_2 , T_4 , T_7 , T_9 and T_{14} are independently hydrogen or
hydroxy protecting group.

42. The taxane of claim 40 wherein R_7 is acyloxy.

43. A taxane having the structure:



wherein

M is a metal or comprises ammonium:

R₁ is hydrogen, hydroxy, protected hydroxy, or together with R₁₄ or R₂ forms a carbonate;

R₂ is keto, -OT₂, acyloxy, or together with R₁ forms a carbonate;

R₄ is -OT₄ or acyloxy;

R₇ is acyloxy;

R₉ is hydrogen, keto, -OT₉, or acyloxy;

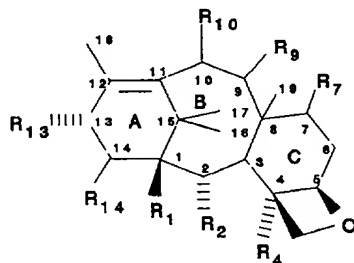
R₁₀ is hydrogen, keto, -OT₁₀, or acyloxy;

R₁₃ is hydroxy, protected hydroxy, keto, or MO-;

R₁₄ is hydrogen, -OT₁₄, acyloxy or together with R₁ forms a carbonate; and

T₂, T₄, T₉, T₁₀ and T₁₄ are independently hydrogen or hydroxy protecting group.

44. A taxane having the structure:



wherein

M is a metal or comprises ammonium:

R_1 is hydrogen, hydroxy, protected hydroxy, or together with R_{14} or R_2 forms a carbonate;

R_2 is keto, $-OT_2$, acyloxy, or together with R_1 forms a carbonate;

R_4 is $-OT_4$ or acyloxy;

R_7 is $-OT_7$;

R_9 is hydrogen, keto, $-OT_9$, or acyloxy;

R_{10} is hydrogen, keto, $-OT_{10}$, or acyloxy;

R_{13} is hydroxy, protected hydroxy, keto, or $MO-$;

R_{14} is hydrogen, $-OT_{14}$, acyloxy, or together with R_1 forms a carbonate;

T_7 is a ketal or an acetal radical;

T_2 , T_4 , T_9 , T_{10} , and T_{14} are independently hydrogen or hydroxy protecting group.

45. The taxane of claim 44 wherein

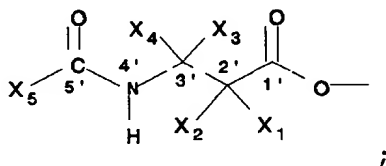
R_1 is hydroxy or together with R_{14} or R_2 forms a carbonate;

R_2 is $-OCOZ_2$, $-OCOOZ_2$, or together with R_1 forms a carbonate;

R_4 is $-OCOZ_4$;

R_9 is hydrogen or keto;

R_{13} is hydroxy, protected hydroxy, or



R_{14} is hydrogen, hydroxy, protected hydroxy, or together with R_1 forms a carbonate;

X_1 is $-OX_6$ or $-NX_8X_9$;

X_2 is hydrogen, hydrocarbyl, or substituted hydrocarbyl;

- 15 X₃ and X₄ are independently hydrogen, hydrocarbyl,
substituted hydrocarbyl, or heteroaryl;

X_5 is $-X_{10}$, $-OX_{10}$, or $-NX_8X_{10}$;

X₆ is a hydroxy protecting group;

X₈ is hydrogen, hydrocarbonyl, or substituted

20 hydrocarbyl;

X₉ is an amino protecting group;

X₁₀ is hydrocarbyl, substituted hydrocarbyl, or heteroaryl; and

Z₂ and Z₄ are independently hydrocarbyl, substituted

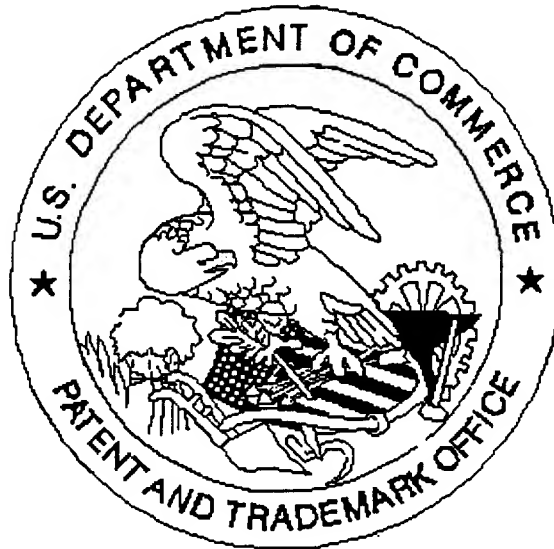
25 hydrocarbonyl, or heteroaryl.

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